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UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new nonprovisional applications under 37 C.F.R. § 1.53(b))

Attorney Docket No. 75329 77432

First Inventor or Application Identifier Thomas J. Shaw

Title Syringe Plunger Assembly and Barrel

Express Mail Label No. EL465447941US

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

1. * Fee Transmittal Form (e.g., PTO/SB/17)
(Submit an original and a duplicate for fee processing)
2. Specification [Total Pages 46]
(preferred arrangement set forth below)
 - Descriptive title of the Invention
 - Cross References to Related Applications
 - Statement Regarding Fed sponsored R & D
 - Reference to Microfiche Appendix
 - Background of the Invention
 - Brief Summary of the Invention
 - Brief Description of the Drawings (if filed)
 - Detailed Description
 - Claim(s)
 - Abstract of the Disclosure
3. Drawing(s) (35 U.S.C. 113) [Total Sheets 8]
4. Oath or Declaration [Total Pages 2]
 - a. Newly executed (original or copy)
 - b. Copy from a prior application (37 C.F.R. § 1.63(d))
(for continuation/divisional with Box 16 completed)
 - i. DELETION OF INVENTOR(S)
 - Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §§ 1.63(d)(2) and 1.33(b).

* NOTE FOR ITEMS 1 & 13 IN ORDER TO BE ENTITLED TO PAY SMALL ENTITY FEES, A SMALL ENTITY STATEMENT IS REQUIRED (37 C.F.R. § 1.27), EXCEPT IF ONE FILED IN A PRIOR APPLICATION IS RELIED UPON (37 C.F.R. § 1.28).

16. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below and in a preliminary amendment:

 Continuation Divisional Continuation-in-part (CIP)

of prior application No: 08/843,050

Prior application information: Examiner Yasko, J.

Group / Art Unit: 3734

For CONTINUATION or DIVISIONAL APPS only: The entire disclosure of the prior application, from which an oath or declaration is supplied under Box 4b, is considered a part of the disclosure of the accompanying continuation or divisional application and is hereby incorporated by reference. The incorporation can only be relied upon when a portion has been inadvertently omitted from the submitted application parts.

17. CORRESPONDENCE ADDRESS

<input checked="" type="checkbox"/> Customer Number or Bar Code Label		20873 (Insert Customer No. or Attach bar code label here)		or <input type="checkbox"/> Correspondence address below	
Name	Harry J. Watson				
Locke Liddell & Sapp LLP					
Address	2200 Ross Ave., Ste. 2200				
City	Dallas	State	TX	Zip Code	75201
Country		Telephone	214 740 8713	Fax	214 740 8800

Name (Print/Type)	Harry J. Watson	Registration No. (Attorney/Agent)	29,985
Signature	<i>Watson</i>		
	Date 7/17/2000		

Burden Hour Statement: This form is estimated to take 0.2 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Box Patent Application, Washington, DC 20231.

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: Shaw, Thomas J.

Application No.:

Group No.:

Filed:

Examiner:

For: Syringe Plunger Assembly and Barrel

**Assistant Commissioner for Patents
Washington, D.C. 20231**

EXPRESS MAIL CERTIFICATE

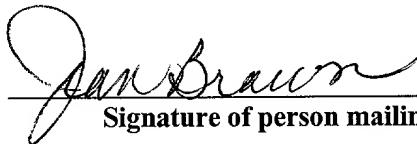
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I hereby state that the following *attached* paper or fee

Utility Patent Application Transmittal, Fee Transmittal, Preliminary Amendment, prior filed application with drawings, declarations and power of attorney, Express Mail Certificate of Mailing

is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 C.F.R. section 1.10, on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231.

Jean Brown



Signature of person mailing paper or fee

**DECLARATION CLAIMING SMALL ENTITY
STATUS PURSUANT TO 37 CFR 1.9(f) and 1.27 (b)**

SOLE INDEPENDENT INVENTOR

As a below named inventor, I hereby declare that:

I qualify as an independent inventor as defined in 37 CFR 1.9(c) for purposes of paying reduced fees under section 41(a) and (b) of Title 35, United States Code, to the Patent and Trademark Office with regard to the invention entitled:

"SYRINGE PLUNGER ASSEMBLY AND BARREL"

described in the specification filed herewith.

Each person, concern or organization to which I have assigned, granted, conveyed, or licensed or am under an obligation under contract or law to assign, grant, convey, or license any rights in the invention is listed below:

- [] no such person, concern, or organization.
[X] persons, concerns or organizations listed below*

Retractable Technologies, Inc.

*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b)).

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Name and address of inventor: Thomas J. Shaw
1510 Hillcrest
Little Elm, Texas 75068

Signature

4/27/97
Date

69609

**DECLARATION CLAIMING SMALL ENTITY STATUS
PURSUANT TO 37 CFR 1.9(f) and 1.27(c)**

SMALL BUSINESS CONCERN

I hereby declare that I am an officer of the small business concern empowered to act on behalf of the concern identified below:

NAME OF CONCERN:	Retractable Technologies, Inc.
ADDRESS OF CONCERN:	511 Lobo Lane P. O. Box 9 Little Elm, Texas 75068

The above identified small concern qualifies as a small business concern as defined in 13 CFR 121.3-18, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees under §41(a) and (b) of Title 35, United States Code, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is averaged over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year, and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.

Rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention, entitled:

SYRINGE PLUNGER ASSEMBLY AND BARREL

by inventor: Thomas J. Shaw

described in the specification filed herewith.

Rights held by the above identified small business concern are not exclusive, each individual, concern or organization having rights to the invention is listed below*

Thomas J. Shaw

*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities (37 CFR 1.27)

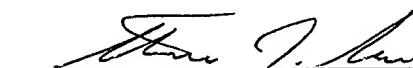
No rights to the invention are held by any person, other than the inventor, who could not qualify as a small business concern under 37 CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e).

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. [37 CFR 1.28(b)]

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Name and Title: Thomas J. Shaw
President

Address: 511 Lobo Lane
P.O. Box 9
Little Elm, Texas 75068



Signature

4/1/97
Date

75329 77432

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PATENT APPLICATION

Serial Number:

Filing Date:

Applicant: Thomas J. Shaw

Title: Syringe Plunger Assembly and Barrel

Group Art Unit:

Examiner:

Assistant Commissioner of Patents
Washington, D. C. 20231

Sir:

PRELIMINARY AMENDMENT

In the Title

5 Please amend the title to read as follows:

-- A Retractable Syringe Assembly Designed for One Use. --

In the Cross Reference to Related Application

Please amend the Cross Reference to Related Applications to read as follows:

-- This is a continuation of copending patent application serial number 08/843,050 filed

10 April 25, 1997 entitled "Syringe Plunger Handle Assembly and Barrel which was a continuation-in-part of patent application serial number 08/537,242 filed September 29, 1995 entitled "Tamperproof Retractable Syringe", now U.S. Patent 5,632,733 which in turn was a continuation-in-part of patent application serial number 08/438,954 filed May 11, 1995 now U.S. Patent 5,578,011, all by the same inventor for which benefit of 35 U.S.C. § 120 is claimed. --

In the Claims

The present application as filed contains claims 1 - 28. Before calculating the filing fee, please substitute new claims 29 - 55 for the existing claims.

29. A retractable syringe assembly designed for one use, comprising:

5 a hollow syringe body comprising a barrel having a front end portion containing a retraction mechanism having a retractable needle and continuous retaining member configured for operation by forward movement of a plunger;

the continuous retaining member having one outside mating surface making a seal for a variable fluid chamber in the barrel;

10 a plunger having a front end portion comprising a head and a supporting surface on the front end portion having a plunger seal element fixed on the supporting surface;

the plunger being reciprocally mounted in said barrel with the plunger seal element in sliding sealed contact with the barrel; and

15 the retractable needle retraction mechanism being released for retraction when the plunger is moved forward to release the continuous retaining member, without contact between the plunger seal element and the continuous retaining member and without relative movement between the plunger seal element and its supporting surface.

30. The assembly of claim 29 wherein a structure mounted in the front end portion of the barrel prevents forward motion of the retractable needle during retraction of the needle to prevent pain when the needle is retracted from a patient.

31. The assembly of claim 29 wherein the plunger carries a part which protrudes to contact the continuous retaining member and release the retractable needle when retraction is initiated by pushing on the plunger.

32. The assembly of claim 29 wherein the continuous retaining member is a separable part of the retraction mechanism which acts as a fluid seal for a variable chamber in the barrel behind the separable part.

33. The assembly of claim 31 wherein the continuous retaining member is separable from the retractable needle when retraction is initiated by pushing on the plunger.

34. The assembly of claim 33 wherein the continuous retaining member is separated from the retractable needle by means of force applied by said part to said continuous retaining member when retraction is initiated by pushing on said plunger.

35. The assembly of claim 29 wherein the plunger has an end cap and the hollow syringe body has a back end portion having an opening for receiving the end cap.

36. The assembly of claim 35 wherein the plunger end cap is lodged in the opening of the back end portion of the hollow syringe body by pressing the end cap to cause retraction, thereby stopping the plunger from further forward movement and preventing reuse.

37. A retractable syringe assembly designed for one use, comprising:

a hollow syringe body comprising a barrel having a front end portion in which a retraction mechanism is mounted, the retraction mechanism having a needle which is retractable and a continuous retaining member which holds the retractable needle;

5 the retraction mechanism being operable by forward movement of a plunger without distorting the barrel;

a plunger having a front end portion comprising a head and a supporting surface located on the front end portion, with a plunger seal element fixed on the supporting surface;

the plunger being reciprocally mounted in said barrel with the plunger seal element in sliding sealed contact with the barrel; and

5 forward movement of the plunger releases the needle from the continuous retaining member by applying a separating force to the continuous retaining member, without the aid of the plunger seal element and without relative movement between the plunger seal element and its supporting surface.

38. The assembly of claim 37 wherein the continuous retaining member acts as a fixed seal for a variable chamber in the barrel behind the separable part.

39. The assembly of claim 38 wherein a structure mounted in the front end portion of the barrel prevents forward motion of the retractable needle during retraction of the needle to prevent pain when the needle is retracted from a patient.

40. The assembly of claim 37 wherein the continuous retaining member is separable from the retractable needle when retraction is initiated by pushing the plunger to move it forward with respect to the barrel.

41. The assembly of claim 40 wherein the plunger carries a part which protrudes to contact the continuous retaining member and release the retractable needle when retraction is initiated by pushing on the plunger.

42. The assembly of claim 41 wherein the continuous retaining member is separated
10 from the retractable needle by means of force applied by said part to said continuous retaining member when retraction is initiated by pushing on said plunger.

43. The assembly of claim 37 wherein the plunger has an end cap and the hollow syringe body has a back end portion having an opening for receiving the end cap.

44. The assembly of claim 43 wherein the plunger end cap is lodged in the opening of the back end portion of the hollow syringe body by pressing the end cap to cause retraction, thereby stopping the plunger from further forward movement and preventing reuse.

45. A retractable syringe assembly designed for one use, comprising:

a hollow syringe body comprising a barrel having a front end portion in which a retraction mechanism is mounted, the retraction mechanism having a needle which is retractable and a continuous retaining member which holds the needle;

5 the continuous retaining member having one outside mating surface making a seal for a variable fluid chamber in the barrel;

a plunger having a front end portion comprising a head and a supporting surface located on the front end portion, with a plunger seal element fixed on the supporting surface;

10 the plunger being reciprocally mounted in said barrel with the plunger seal element in sliding sealed contact with the barrel; and

the retractable needle retraction mechanism being released for retraction when the plunger is moved forward to release the continuous retaining member, without the plunger seal element going beyond said one outside mating surface and without motion of the plunger seal element relative to its supporting surface.

46. The assembly of claim 45 wherein the continuous retaining member acts as a fixed seal for a variable chamber in the barrel behind the separable part.

47. The assembly of claim 46 wherein a structure mounted in the front end portion of the barrel prevents forward motion of the retractable needle during retraction of the needle to prevent pain when the needle is retracted from a patient.

48. The assembly of claim 45 wherein the continuous retaining member is separable from the retractable needle when retraction is initiated by pushing the plunger to move it forward with respect to the barrel.

49. The assembly of claim 48 wherein the plunger carries a part which protrudes to contact the continuous retaining member and release the retractable needle when retraction is initiated by pushing on the plunger.

50. The assembly of claim 49 wherein the continuous retaining member is separated from the retractable needle by means of force applied by said part to said continuous retaining member when retraction is initiated by pushing on said plunger.

51. The assembly of claim 45 wherein the plunger has an end cap and the hollow syringe body has a back end portion having an opening for receiving the end cap.

52. The assembly of claim 51 wherein the plunger end cap is lodged in the opening of the back end portion of the hollow syringe body by pressing the end cap to cause retraction, thereby stopping the plunger from further forward movement and preventing reuse.

53. A retractable syringe assembly designed for one use, comprising:

a hollow syringe body comprising a barrel having a front end portion containing a retraction mechanism having a retractable needle and continuous retaining member configured for operation by forward movement of a plunger;

a biasing element mounted in the front of the barrel;
a plunger having a front end portion comprising a head and a supporting surface on the front end portion having a plunger seal element fixed on the supporting surface; and
where the front end of the barrel has an inwardly sloping wall continuously sloping to the
5 point of contact with the continuous retaining member.

54. A retractable syringe assembly designed for one use, comprising:

a front end retraction mechanism and retractable needle including a continuous retaining member;

a plunger having a front end portion comprising a head and a supporting surface on the
10 front end portion having a plunger seal element fixed on the supporting surface;

a rigid plunger seal element stop surface which acts as a plunger seal element stop; and

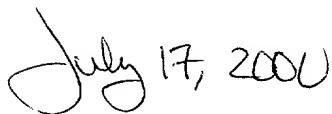
where the retraction mechanism is operated by forward movement of the plunger to release the retractable needle for retraction while the plunger seal element remains fixed to its supporting surface.

55. The assembly of claim 54 wherein the plunger operates the retraction mechanism by acting on the continuous retaining member to release the retractable needle for retraction while the plunger seal element remains fixed to its supporting surface.

Respectfully submitted,



Harry J. Watson
Registration No. 29,985



LOCKE LIDDELL & SAPP LLP
2200 Ross Avenue, Suite 2200
Dallas, Texas 75201
214/740-8000 Telephone
214/740-8800 Facsimile

SYRINGE PLUNGER ASSEMBLY AND BARREL**CROSS REFERENCE TO RELATED APPLICATION**

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This is a continuation-in-part of copending patent application serial number 08/537,242 filed September 29, 1995 entitled Tamperproof Retractable Syringe which in turn was a continuation of serial number 08/438,954 filed May 11, 1995, now U.S. Patent No. 5,578,011 all by the same inventor for which benefit is claimed under 35 U.S.C. §120.

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FIELD OF THE INVENTION

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This invention relates to a medical device, and more particularly to a retractable syringe and components suitable for mass production and assembly having a low triggering force and high blowout pressure which is nonreusable after one use.

BACKGROUND OF THE ART

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A major cause to the spread of AIDS in the general population is the presence of IV drug users who share and reuse hypodermic syringes to inject drugs. Infection can be spread from AIDS patients in hospitals and medical facilities through accidental needle sticks from needles used on infected patients. Used syringes with extended needles present a risk to medical personnel and sanitation employees and others in the disposal chain.

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The gravity of the threat posed by AIDS and the fact that the main vector of the spread of the dreaded disease is through reuse of syringes by IV drug users has resulted in intense activity to develop the most practical, most reliable, easily assemblable, mass-producible syringe.

There are a number of syringes of different designs which have needles which will retract at the end of the injection cycle. Most of these have never reached the market because of

various deficiencies. Prime among the usual deficiencies of the prior art are problems of complexity, reliability, cost and ease of use. The most commonly used syringes are 1cc and 3cc syringes which must be mass-produced at the rate of millions per day. Cost is a significant factor both in manufacture of the parts and assembly of the device. High speed production requires molds with 64 cavities or more to reduce unit cycle time. Therefore, molded structures within the barrel that require collapsing core pins such as are shown in much of the art are unlikely to be producible at competitive costs.

One of the problems of the prior art of retractable syringes is the sheer number and complexity of parts which must be formed and assembled. Other problems with the prior art are dependence on flexing or breaking of internal parts by the plunger in order to release the retraction mechanism and use of a diaphragm at the end of the plunger which must be penetrated by a needle holding member and spring. These structures present serious quality control and assembly problems. Small broken off pieces can present a risk of hang-ups. Hooks are often used to releaseably secure retraction mechanisms. Hooks present difficult holding and control problems, may cause retention of air bubbles upon filling and may be undesirably temperature sensitive.

The prior art frequently has a two-piece barrel in order to be able to assemble a retraction device in the nose. This requires at least an additional part and assembly step. It is still necessary to pass the sharp injection needle through a small opening often while compressing a spring before the two parts can be assembled. The tiny needles are produced in the form of coil tubing and vary significantly from straightness after they are cut to length. This leads to

difficult assembly problems if the needle must be passed through a small opening. The extremely sharp tip will catch the edge of a hole and jam the production line.

The rare prior art that employs a front mounted retraction mechanism in a one-piece barrel with a plugged hollow plunger, Tsao U.S. Pat. No. 5,084,018, among other things does 5 not show reduced barrel area to prevent excessive blowout pressure, employs engaging flanges to secure all retraction parts, requires concurrent distortion of internal parts and flanges to effect release, cumulating in excessive force required to retract and requires ventilation holes because of a compartmented barrel.

The prior art has not produced a retractable nonreusable tamperproof syringe for mass 10 production and assembly which is simple, reliable, cost effective, easy to use and retract, looks like a conventional syringe, has few parts which are easy to make and assemble, is not temperature sensitive and not subject to danger of premature retraction.

The prior art has not recognized a retraction mechanism with separable parts that relies 15 entirely on clamping force or friction at a smooth walled reduced diameter transition zone in the barrel with mating lands which are slidably or separably released in response to relatively low thumb pressure while having resistance to premature retraction and high blowout pressure resulting from high pressure produced in the fluid chamber during an injection. The prior art has not recognized that such a structure can be molded as a one piece outer body over a core that can be pulled out from behind allowing the retraction mechanism to be easily pushed into 20 place from behind, steered by the narrow nose portion. Neither does the prior art in such a combination realize the desirable non-cumulation of forces resisting retraction in order to

minimize the thumb force required, having a most simple tamperproof feature and the fewest number of easily made parts.

The syringe plunger assembly has a combination of features not found in a prior art syringe. A head end which acts like a piston when installed in a syringe barrel has a reduced 5 diameter front end having an opening and a dislodgeable stopper slidingly mounted in the opening projecting forwardly from the tip. Cooperating lands within the opening and on the head of the dislodgeable stopper seal the opening into the hollow interior of the plunger. The area of the stopper is relatively small when compared to the area exposed to the piston, which compresses fluid in a chamber below the piston. The ratio of the total area of the fluid chamber to the fluid exposed area of the stopper is at least two to one, more preferably three to one or 10 more so that the stopper requires less holding force without blowing out back into the internal cavity. The cooperating lands have sufficient length so that the stopper can move back to the tip when the plunger moves forward at the end of an injection stroke without unsealing the plunger opening. A reduced holding force is sufficient to prevent blowout of the stopper after 15 the stopper has been moved back to the tip because the stopper is exposed to a lower pressure generated force because of its relatively smaller area. The back of the plunger is vented so that entry of retractable parts which upon retraction finish dislodging the stopper and carry it back into the cavity, do not generate internal pressure that can blow out the nose of the syringe carrying any residual fluid with it. The thumb cap on the plunger is received and recessed into 20 the opening at the back of the barrel when retraction occurs. The plunger cannot be grasped after this occurs to help prevent reuse.

These features and more are found in the inventive combination herein further disclosed which is especially suited for high speed production and assembly at low cost.

SUMMARY OF THE INVENTION

The invention is a reliable retractable tamperproof syringe having multiple tamperproof features which operates on a principle which permits low cost parts which are few in number and well suited for high speed mass production and assembly. The syringe structure features a one piece hollow outer body having a longitudinally extending wall which is stepped. The wall comprises an elongated barrel and nose with a transition zone connecting the barrel and nose. The nose has a reduced diameter relative to the barrel. The outer body has an inwardly facing surface in the wall at the most constricted part of the transition zone where the nose begins. A plunger assembly is disposed partially within the elongated barrel with an end cap for depression of the plunger extending from an opening in the back of the barrel. The head of the plunger, which has a retraction cavity for receiving parts of a retraction mechanism, moves in slidable sealed contact with the interior of the barrel.

A retraction mechanism is lodged in the nose of the body. The retraction mechanism comprises an elongated needle holder and spring combination wherein the needle holder has an elongated body with a needle holding portion in front and a head in back. The head of the needle holder has a cooperating outwardly facing surface configured to cooperate with said inwardly facing surface along an interface oriented in the direction of retraction to produce a holding force on the needle holder when installed in the nose in the unretracted position. The needle holder and spring are easily installable from the rear of the barrel toward the nose and

releaseably held by sliding engagement of said cooperating inwardly and outwardly facing surfaces while compressing the spring and thereby producing a holding force on the needle holder in opposition to the retraction force applied to the needle holder by the spring. The parts are circular in cross section.

5 The outwardly facing surface on the circular head of the needle holder is slightly greater in diameter than the circular inward facing surface in the wall at the most constricted portion where the nose begins. The needle holder is thus clamped in position by hoop stresses induced in the outer body and held in position by frictional holding force. The needle holder is released in response to depression of the plunger to a retraction position. Retraction occurs in response to thumb force on the plunger when a portion of the plunger passing into the transition zone separates at least a portion of the inwardly and outwardly facing cooperating surfaces thereby reducing the holding force on the needle holder to an amount less than a retraction force on the needle holder produced by the spring whereby the needle holder is retracted into the cavity a distance sufficient to withdraw an injection needle, attached to the needle holder, into the outer body.

10 In one embodiment, the head of the needle holder is a two part head comprising an inner head surrounded by a separable retainer member wherein the outer surface of the retainer member is the outwardly facing surface with cooperates with the inwardly facing surface in the wall to retain the needle holder in an unretracted position at the most constricted part of the transition zone where the nose begins. The retainer member is a ring member coupled to the inner head along a sliding interface oriented in the direction of retraction with a friction force which exceeds the retraction force provided by the spring. The front of the needle holder is

grounded in the nose portion against forward movement. The plunger head is configured to pass through the most constricted area and push against the retainer member without also pushing against the head of the needle holder. An alternate construction of the two part head of the needle holder comprises the separable retainer member being tack welded to the inner head of
5 the needle holder, preferably along a very small ridge or bridge between the mating surfaces which holds the two part head together until the bridge is ruptured by movement of the plunger after an injection has occurred.

The front of the plunger has an opening for a stopper slidingly fitted therein in an interference fit. The stopper is fitted in the opening in an interference fit along a sliding interface oriented in the direction of retraction. The stopper is mostly or fully dislodged by contact with the retraction mechanism at the end of an injection cycle by continued depression of the plunger from a first position at the end of the injection cycle to a second position with the tip of the plunger in contact with the retainer ring. This avoids cumulation of the force on the plunger required to dislodge the stopper from the opening and the force required to dislodge the retainer member from the head of the needle holder and outer body wall. Upon further depression of the plunger from the second position to the retraction position, the frictional holding force on the needle holder is reduced until the retraction force provided by the spring exceeds the remaining holding force and the needle holder and needle connected thereto are ejected into the cavity carrying the dislodged stopper along with them. The dislodging of the
10 stopper and the retainer member alone make the syringe non-reusable. The plunger cannot be removed after retraction because the graspable end cap enters an opening at the back of the
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barrel when the plunger is depressed to the retraction position to prevent tampering after retraction.

The retraction cavity of the plunger is preferably vented to prevent a puff of air coming forward at the instant of retraction from blowing a tiny amount of retained fluid from the nose.

5 This condition can occur if the plunger is fully depressed to release the needle holder and dislodge the stopper while the needle is physically restrained from retracting by the septum of a vial which has just been filled with fluid from the syringe. The thumb cap at the rear of the syringe is preferably provided with channels in fluid communication with the interior in cooperation with a closure removably installed in a centrally located opening in the thumb cap.

10 One or more stepped portions of the opening and closure provide seating for the closure. Undercut portions at the side of the closure together with grooves in the interior surface of the plunger wall create passages for air to vent through channels on the thumb cap. This structure prevents air from being trapped by the user's thumb when the thumb cap is pressed to fire the syringe. One or more slots at the back of the barrel around the opening which receives the thumb cap prevent vented air from being trapped by the user's thumb when the plunger is fully depressed.

15 The syringe has a high blowout pressure and a low plunger thumb force required to cause retraction. Blowout pressure is the fluid pressure operating on the stopper and retainer ring during an actual injection. High blowout pressure resistance is obtained because the retainer ring is mounted in the most constricted portion of the barrel where the nose begins which significantly reduces the amount of area exposed to fluid pressure. The smaller retainer ring allows the use of a small needle holder such that the opening in the plunger and the stopper can

be only a fraction of the cross sectional area of the fluid chamber below the plunger head. The ratio of the greatest cross sectional area of the variable chamber and that of the dislodgeable stopper or the ring member are selected so that the maximum expected thumb force on the plunger during an injection will produce a maximum pressure in the chamber which will generate a blowout force on the stopper and retainer member slightly less than the amount of dislodging force necessary to dislodge the stopper and retainer member during retraction. This ratio should be at least two to one, or more preferably three to one or more, in order to ensure against premature blowout of the stopper or retainer ring.

In an alternate embodiment, the fewest number of easily made separate parts are used in a retractable syringe. The alternate embodiment has a similar stopper in the head of the plunger and a similar needle holder and spring combination with mating cooperating inwardly facing and outwardly facing interengaged surfaces at the most constricted part of a transition zone where the nose begins. In the alternate embodiment, there is no retainer ring around the head of the needle holder. Instead a tiny ramp is provided at the transition zone or adjacent the transition zone whereby the head of the plunger gently spreads the barrel outwardly while dislodging the stopper thereby reducing the clamping or friction force on the head of the needle holder provided by the wall of the outer body. The holding force is thereby reduced below the retraction force provided by the compressed spring and the needle holder is ejected into the cavity of the plunger carrying the dislodged stopper along with it.

Manufacture and assembly is facilitated by the fact that the plunger and the outer body can be molded with a non-collapsible core tool that can be pulled out from behind. The parts are simply shaped and do not have hooks and parts with reentrant angles that require collapsible

core pin technology. The outer body can be made in one piece and assembled from the rear. The narrowed nose portion provides no lateral space which will permit bunching of the spring and jamming when the retraction assembly is moved forward in the outer body. In fact, the nose serves as a guide to steer the parts into the proper position in one smooth stroke.

The needle does not have to be installed before the retraction mechanism is put in place because it is readily installed from the front after the needle holder is slidingly lodged in the nose. Significant variations in the holding force on the needle holder and the dislodging force on the stopper due to slight variances in the tolerance of the mating parts is avoided because the longitudinal wall of the outer body has some flexibility. The wall can spread outwardly slightly and the stopper and head of the needle holder can compress slightly radially and expand slightly in the longitudinal direction to avoid significant changes in the holding force caused by small changes in the actual diameters. Consistency in the amount of retraction force is thereby provided and economy is assured.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a cross section along the central axis of a first embodiment of the invention with the plunger positioned in a first position at the end of an injection cycle;

Figure 2 is the syringe of Figure 1 with the plunger depressed additionally to dislodge the stopper at a second position of the plunger wherein the tip of the plunger is ready to operate the retraction mechanism;

Figure 3 is the syringe of Figure 2 wherein the plunger has been further depressed to a retraction position, retraction has occurred and the cap at the back of the plunger is closely received in an opening at the back of the outer body;

Figure 4A is a partial cross section on the central axis of an alternate tamperproof opening in the back of the outer body prior to retraction;

Figure 4B is the structure of Figure 4A with the plunger in the retracted position received in an opening at the back of the outer body;

Figure 5 is a cross section along the central axis of a simplified alternate syringe structure without a retainer member around the needle holder, which is released by separation of the friction surfaces, shown in the plunger position which represents the end of an injection cycle;

Figure 6 is the syringe structure of Figure 5 wherein the plunger is further depressed to dislodge the stopper and begin to release the friction surfaces just prior to retraction;

Figure 7 is the syringe structure of Figure 6 with the plunger further depressed beyond the position of Figure 6 to the retraction position where retraction has occurred and the cap is secure within an opening in the back of the hollow outer body.

Figure 8 is a schematic longitudinal cutaway view in elevation through the center of the two part head showing how a tack weld can be applied to simultaneously seal and hold the retainer ring in place on the needle holder.

Figure 9 is an exploded perspective view showing the barrel and retraction mechanism of Figure 1 with a modified plunger assembly;

Figure 10 is a plan view of the thumb cap of the plunger assembly shown in Figure 9 with the preferred closure;

Figure 11 is a cut away elevational view of the structure at the back end of the plunger and end cap of Figure 9 and 10 along line 11-11 showing the preferred closure;

Figure 12 is a cut away elevational view of the plunger end cap and closure of Figure 11 as the thumb cap is just being received into the barrel opening;

5 Figure 13 is a plan view of a first alternative thumb cap and closure combination utilizing a flat sided closure and four channels in the thumb cap;

Figure 14 is a cut away elevational view on the lines 14-14 of the thumb cap closure combination of Figure 13;

10 Figure 15 is a plan view of a second alternate thumb cap and closure combination with four channels in the thumb cap and undercut portions to provide a vent passage;

15 Figure 16 is a cut away elevational view on the lines 16-16 of the combination of Figure 16.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

In the description that follows, like parts will be referred to by the same reference numerals. Parts with a subscript letter are mean to illustrate a minor variation of a part with the same number. The drawings are enlarged significantly in order to show the details of the invention but generally reflect the true scale which is contemplated. The parts as shown are understood to be preferably circular and symmetrical as is conventional for syringes. The drawings reflect a syringe structure typically having a 1cc to 3cc injection fluid capacity.

20 Figure 1 shows the structure of the first embodiment generally referred to by reference numeral 10. Syringe 10 has a one piece hollow outer body 12. Body 12 has a longitudinally extending wall comprising an elongated barrel 14 and a nose 16 with a transition zone 18

connecting the barrel and nose. A front mounted retraction mechanism lodged in the nose is generally referred to by the reference numeral 20. It comprises the combination of an elongated needle holder 22 and spring 24. The needle holder has an elongated body with a needle holding portion 26 in front for holding a needle 28 and a head 30 in back. Head 30 may consist of a two part head as in Figures 1 - 3 or a one part head as in Figures 5 - 7. The needle holder is released by depression of a plunger that will be described.

A plunger generally designated by the reference numeral 32 is disposed for use partially within barrel 14. The plunger has a head and seal generally referred to by reference numeral 34, in slideable sealed contact with the interior of barrel 14 of outer body 12. The plunger has a seal element 36 that is conventional and a retraction cavity 38 therein.

Head 34 has a tip portion 40 forming an opening 41 into retraction cavity 38. A resilient dislodgable stopper 42 is sealingly positioned in opening 41 with a front portion thereof extending beyond tip 40. Head portion 34 and the back part of stopper 42 have cooperating lands 44, 46, respectively, which seal opening 41. Plunger 32 has an end cap 48 for depression of the plunger by the thumb. End cap 48 has a central opening for permanently receiving force fit plug 50 to close retraction cavity 38 at the back end.

A plurality of longitudinally extending flutes 52 slidably support plunger 32 in barrel 14. In the embodiment of Figure 1, outer body 12 has a collar 54 extending behind finger grips 56 having opening 58 which closely receives the outer periphery 60 of cap 48 when the plunger is depressed to the retracted position. An alternate arrangement is shown in Figures 4A and 4B in which barrel 14 is extended longitudinally, if necessary, so that end cap 48 fits closely within an opening at the back of the barrel where the finger grips are. Figure 4B shows the

tamperproof position with the plunger in the retracted position. It should be noted that depending on the relationship of the inside diameter of the barrel and the diameter of the end cap, the end cap could instead be received right inside the opening at the back of the barrel .
5 Regardless of how the end cap in back of the outer body and barrel are configured, the plunger can no longer be grasped after retraction has occurred because end cap 48 is depressed into an opening.

The wall of outer body 12 and head 30 of the needle holder have mating cooperating smooth surfaces which hold needle holder 22 in the position shown in Figure 1 with spring 24 compressed. Nose 16 has a reduced diameter relative to the barrel. The outer body has a most constricted part where head 30 of needle holder 22 is engaged and held. The outer body has an inwardly facing surface 62 at the most constricted part of the transition zone where nose 16 begins. Similarly, head 30 has an outwardly facing surface 64 configured to cooperate with inwardly facing surface 62 to produce a holding force on needle holder 22 when the retraction mechanism is installed in the nose from the rear. Mating surfaces 62, 64 constitute a sliding interface oriented in the direction of retraction, which seals nose 16. Mating surfaces 62, 64 are preferably friction surfaces which have an interference sliding fit to apply a frictional holding force which holds needle holder 22 in position by friction between the mating parts. It is within contemplation of the invention that one or more of the cooperating interface surfaces could employ a coating or adhesive bond which is ruptured or released when the mating surfaces or
20 lands are separated or moved relative to each other.

Head 30 provides a lower boundary for a variable fluid chamber 68 below head 34. Needle holder 22 has a fluid path 70 in fluid communication with fluid chamber 68 and needle

28. Needle holder 22 has a smaller diameter inner head 72 which is part of head 30. Retainer member 66 is coupled to inner head 72 along sliding interface 74 oriented in the direction of retraction. Retainer member 66 is coupled to inner head 72 with a holding force which exceeds a retraction force applied to the underside of inner head 72 by means of the end of compressed spring 24. A reduced diameter portion 27 of needle holder 22 protrudes through an opening in front 76 of nose 16.

Importantly, retainer member 66 can be visualized as an annular ring surrounding circular inner head 72. The location of retainer member 66 at the most constricted part of the transition zone where the nose begins and the relatively small area exposed to pressurized fluid in chamber 68 results in a high blowout pressure. Since the front portion 26 of the needle holder is grounded or bottomed inside front 76 of nose 16, no amount of pressure will allow needle holder 22 or needle 28 to move forward. Blowout pressure may be defined as the pressure in chamber 68 acting on the exposed area of retainer member 66 to produce a force sufficient to overcome the holding force such that retainer 66 could "blowout" by moving forward and prematurely release needle holder 22.

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Some users have strong hands and might, at the outer limit in an emergency, be able to generate a force of as much as fifteen to eighteen pounds on the plunger during an injection. It is considered almost impossible for anyone to exert a force of more than eighteen pounds. This may be regarded as the maximum expected force which must be taken into account so that ring member 66 will not blowout while an injection is being made. The greatest cross sectional area of variable chamber 68 and the area of retainer member 66 exposed to fluid pressure are selected so that the blowout pressure is higher than the maximum pressure in chamber 68

expected to result from the maximum expected thumb force applied to cap 48 during an injection. This ratio is preferably about two to one and more preferably about three to one or more so that the holding force holding the retraction mechanism in place can be kept at a comfortably low level while the blowout pressure remains high.

5 Dislodgeable stopper 42 has a similar blowout problem to recognize. The front and middle portion of stopper 42 are relieved slightly from opening 41 such that the fluid pressure in chamber 68 is directed against the cross sectional area at cooperating lands 44, 46 and could cause stopper 42 to blowout. A frictional holding force is generated at the lands 44, 46 which may be called a dislodging force which must be overcome to slide stopper 42 rearwardly before retraction. The ratio of the maximum cross sectional area across the interior of variable chamber 68 to the maximum cross sectional area of stopper 42 exposed to pressure in chamber 68 are selected so that the maximum expected thumb force on plunger 32 during an injection will produce a maximum force slightly less than the amount of dislodging force necessary to dislodge the stopper so that stopper 42 will not blowout during an injection. This ratio is preferably not less than about two to one, more preferably three to one or more, whereby a force of about eighteen pounds on the plunger, for example, would produce a pressure generated force of only about nine or six pounds, respectively, on the stopper, so that the stopper can be easily dislodged in advance of retraction at the end of the injection cycle but will not blowout during an injection. The stopper is dislodged after the injection by thumb force applied to the stopper by movement
10 of the plunger.
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The components used for retraction are arranged to avoid cumulation of force during the retraction sequence. In Figure 1, stopper 42 has a forward extension beyond tip 40 which

allows full thumb pressure to be applied to the stopper before any other portion of the retraction mechanism is engaged. The amount of forward extension beyond tip 40 is related to the length of lands 44, 46 such that the forward extension of stopper 42 preferably represents about 80 percent of the engaged land length. When stopper 42 is moved back until the front is even with tip 40, as seen in Figure 2, only about 20 percent of engaged land remains. In Figure 2 it can be seen that thumb force on plunger cap 48 has been applied to partially dislodge stopper 42 such that a gap 78 is created and the remaining engaged land area is represented as area 80.

Since I believe the amount of frictional holding force or dislodging force is roughly proportional to the amount of the length of the sliding interface between cooperating lands 44, 46, it follows, ignoring dynamic effects, that the amount of force remaining decreases as the engaged sliding interface area is reduced. This is what happens as stopper 42 moves back into cavity 38 from the position of Figure 1 to the position of Figure 2. It is believed appropriate to set the initial dislodging force to allow about five pounds at the position of Figure 1 which is reduced to about one pound remaining when the stopper or plug member 42 reaches the position of Figure 2. It might be noted at this point in the description that the front portion of tip 40 preferably has some longitudinally extending slits or openings so that fluid is not trapped in the trapezoidal shaped area of chamber 68, seen in Figure 2, because of contact between tip 40 and the upper surface of retainer ring 66.

Needle holder 22 and spring 24 are combinably installable from the rear of the barrel before the plunger is assembled and releasably held at the most constricted part of the transition zone where the nose begins by sliding engagement of the cooperating inwardly and outwardly facing friction surfaces 62, 64 while compressing spring 24. The length of the engaging land

64 and the amount of interference fit is preferably designed to provide a frictional holding force
in opposition to the retraction force provided by the compressed spring 24 of somewhere around
five pounds even though the spring may apply a retraction force in the retraction direction of
somewhere around a half pound. In use the needle is pushed against a rubber seal in a vial so
5 the needle holder must resist a resulting backward force without being dislodged during the
filling operation. This requirement and blowout pressure limits the low end of the holding force
on the needle holder.

Referring again to Figure 2, it can be seen that further depression of the plunger beyond
the second position of Figure 2 dislodges retainer ring member 66 along the sliding interface 74
provided by the outer surface of inner head 72 and along the inwardly facing friction surface 62.
As the amount of remaining engaged interface is reduced, the amount of force required to
continue moving retainer member 66 off needle holder 22 is reduced and the small remaining
engagement area 80 between lands 44, 46 of the plunger and stopper preferably cause stopper
42 to be dislodged before needle holder 22 is released. When the remaining residual friction
force during continued depression of the plunger becomes less than the retraction force provided
by compressed spring 24, the retraction position of Figure 3 is reached whereby retraction
occurs.

When retraction occurs needle holder 22 moves through opening 41 into cavity 38. The
uncompressed length of spring 24 is selected to provide backward movement sufficient to
20 withdraw an injection needle 28 fixed in front portion 26 entirely within outer body 12, carrying
dislodged stopper 42 with it. At the same time, cap 48 enters opening 58 of the barrel with
peripheral edge 60 closely confined, in order to prevent tampering after retraction. It is

immaterial whether cap 48 moves into the opening at the instant of retraction or after retraction has already occurred because the movement is automatic due to the continued thumb force applied to trigger the retraction. Sufficient unengaged length of inwardly facing friction surface 62 is provided so that retainer member 66 can move downwardly a sufficient distance to reach 5 the retraction position of Figure 3. After retraction, retainer member 66 preferably remains stuck and prevents any possibility of any one being able to reengage it with the head of needle holder 22. The diameter of land 62 in the area designated 63 can be increased slightly to provide relief for retainer ring 66 as it is pushed down by tip 40.

It is also within the contemplation of the invention that separable retainer member 66 may be removably coupled to inner head 72 of needle holder 22 by means of a relatively small in area "tack" weld which is sufficient to resist the retraction force applied to needle holder by spring 24 but which can be ruptured or separated by depression of the plunger beyond the position shown in Figure 2, to release the needle holder and allow retraction. This is schematically illustrated in Figure 8 with respect to alternate head 30a with the parts of syringe body 12 and needle holder 22 cutaway to focus on the modification. The remainder of the syringe structure would be like Figures 1 -3.

In Figure 8, inner head 72a has an outwardly facing surface 74a and a very small raised portion or series of horizontally spaced apart raised portions 73 around the periphery in a continuous band or annular ring which extend relatively uniformly outwardly beyond peripheral 20 surface 74a of head 72a. The raised portion could be on the inner surface 75 of retainer 66a instead of being on surface 74a of the needle holder. The head of the needle holder is preferably

circular but could be conceivably another shape with the retainer member 66a correspondingly configured to conform to it.

The inwardly facing surface 75 of inner head 72a is in contact with raised portion 73 on the outer surface of inner head 72a and there may be a small gap 77 between them all around. 5 The raised portion 73 couples retainer 66a to inner head 72a and may be referred to as a bridging portion which resists the blowout pressure referred to above and holds the needle holder in place against the retraction force imposed on the needle holder by spring 24 together with any small additional forces that may be applied when the needle is pushed against the rubber seal of a vial in preparation for use. The bridging portion may be formed by "tack" welding the raised portion 73 to the inner surface of the ring 66a or by providing any other form of frangible bridging portion that holds the separable ring member 66 and needle holder head 72a together. It is required that however done, the bridging portion must also serve as a seal between the facing surfaces of the ring member and inner head so that fluid under pressure cannot pass from chamber 68 through gap 77 to reach the nose portion of the device. All fluid must pass through fluid passage 70.

It can be seen that when the position of Figure 2 is reached the front tip 40 of the plunger presses against retainer ring 66a after stopper 42 is almost dislodged and uncouples the retainer ring 66a from the inner head 72a of needle holder 22a. Any tack weld connecting the separable parts at the bridging portion is ruptured, fractured or otherwise separated so as to separate 20 retainer ring 66a from inner head 72a thus releasing needle holder 22a from further restraint. They and the force applied by spring 24 causes retraction to occur much as before described and shown in Figure 3.

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It is believed that the increased diameter of the raised portion 73 should be within the range of about 1 to 8 thousandths of an inch which may be dictated by the ability of the molding equipment available to produce a consistent bridging portion without defects. It is believed that it may be desirable to employ different polymeric materials for the retainer ring and needle holder to facilitate tack welding, such as a suitable polyvinyl chloride (PVC) for the retainer ring and a suitable polycarbonate plastic material for the needle holder. One way to couple these two parts may be to assemble them and expose them to a temperature of about 120°C for twenty minutes or so to allow some diffusion or incipient melting to occur where they touch. The raised portion creates a high unit pressure where it comes into contact with the inwardly facing surface of retainer 66a. Sonic welding could also be employed. A coating or adhesive which couples the retainer ring to the needle holder and can be uncoupled by means of force applied to the retainer ring by the plunger is also within the contemplation of the invention.

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An alternate syringe 82 is disclosed in Figures 5 - 7. In Figure 5, Syringe 82 has a one piece hollow outer syringe body 84. Body 84 has a longitudinally extending wall comprising an elongated barrel 86 and a nose 88 with a transition zone 90 connecting the barrel and nose. A front mounted retraction mechanism lodged in nose 88 is generally referred to by the reference numeral 92. It comprises the combination of an elongated needle holder 94 and spring 96. The needle holder has an elongated stem body with a needle holding portion 100 in front for holding needle 28 and a head 102 in back. In this case, head 102 is a one part head integral with the rest of needle holder 94. Spring 96 delivers a retraction force in a retraction direction to the underside of head 102.

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A plunger generally designated by reference numeral 104 is disposed for use partially within barrel 86. Plunger 104 has a head portion 106 which moves in slidable sealed contact with the interior of barrel 86 of outer body 84. Although a separate seal might be used on head 106, this embodiment is suitable for a smaller diameter, such as a 1cc syringe, and can be used with head 106 also serving as the seal. A retraction cavity 108 is provided in the interior of hollow plunger 104. Head 106 has a tip portion 110 forming an opening 112 for a dislodgable stopper 114 having a front portion extending beyond tip 110. Head portion 106 has an inwardly facing land 116 and the back of stopper 114 has an outwardly facing land 118 comprising cooperating friction surfaces which seal opening 112. The back portion of outer body 84 may have finger grips 120 and the same collar 54 and end cap 48 previously disclosed. The alternate arrangement of Figures 4A and 4B may also be employed.

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The outer portion of tip 110 may be equipped with an angled surface 122 designed to cooperate with a small ramp surface 124 located in the vicinity of transition zone 90. The wall of outer body 84 and head 102 of the needle holder have mating cooperating friction surfaces which frictionally hold needle holder 102 in the position shown in Figure 5 with spring 96 compressed. Nose 88 has a reduced diameter relative to barrel 86. The outer body has a most constricted part where the head 102 of needle holder 94 is frictionally engaged. The outer body has an inwardly facing surface or land 126 at the most constricted part of the transition zone where nose 88 begins. Similarly, head 102 has an outwardly facing friction surface 128 configured to cooperate with inwardly facing surface 126 to produce a frictional holding force on needle holder 94 when the retraction mechanism is installed in the nose from the rear.

Mating surfaces 126, 128 constitute a sliding interface oriented in the direction of retraction, which seal nose 88. Mating surfaces 126, 128 are preferably smooth friction surfaces which have an interference sliding fit when needle holder 94 is installed from the rear whereby a frictional holding force holds needle holder 94 in position by friction between land 126 and head 102 of needle holder 94. It is within contemplation of the invention that one or both of these surfaces could have a coating or adhesive bond which is ruptured when the mating surfaces are separated to release the needle holder.

Head 106 provides the upper boundary for a variable fluid chamber 130 below head 106. Needle holder 94 has a fluid path 132 in fluid communication with chamber 130 and needle 28. Needle holder 94 is releasably coupled at surfaces or lands 126, 128 with a holding force that exceed the retraction force applied to the underside of head 102 by the end of compressed spring 96. A reduced diameter portion 134 of needle holder 94 protrudes through an opening in front 136 of nose 88. Blowout pressure is not a factor with respect to the needle holder on the alternate embodiment. No amount of pressure will allow needle holder 94 or needle 28 to move forward since the front portion 100 of the needle holder is grounded or bottomed inside front 136 of nose 88.

Blowout pressure is still a factor to be considered in connection with stopper 114. Blowout pressure would be the pressure in chamber 130 produced by thumb force on cap 48 acting on the cross sectional area of stopper 114 which could overcome the holding force, causing stopper 114 to dislodge from opening 112 prematurely. The ratio of the maximum cross sectional area across the interior of variable chamber 130 to the maximum cross sectional area of stopper 142 exposed to pressure in chamber 130, and the dislodging force necessary to

dislodge stopper 144, are selected so that the maximum expected thumb force on plunger 104 during an injection will not cause the stopper to blowout. Yet the stopper will still be dislodged by the dislodging force on the plunger once the front of stopper 114 contacts the retraction mechanism after the injection has ended. The ratio referred to is preferably not less than about 5 two to one, or more preferably about three to one or more, whereby a force of about eighteen pounds on the plunger, for example, would produce a pressure generated force of only about nine or six pounds respectively, on the stopper, so that the stopper can be easily dislodged in advance of retraction at the end of the injection cycle but will not blowout during an injection. The smaller diameter stopper allows two or three times the thumb force to be used during the 10 injection cycle than required to actually dislodge the stopper by direct application of force.

By reference to Figures 5 - 7, the operation and further features of the alternate embodiment are discussed. The syringe is used in the normal manner until the plunger is depressed to the first position of Figure 5 which is the end of the injection cycle. Stopper 114 has a forwardly extending end which has come into contact with head 102 of needle holder 94 to block fluid path 132. Further depression of plunger 104 toward the position of Figure 6 mostly or fully dislodges stopper 114 and begins spreading barrel 84 at the transition zone by 15 sliding contact between head portion 106 and ramp 124. Ramp 124 is a very small inwardly extending annular thickening of the wall of barrel 86 which can take many shapes or forms. For example, ramp 124 may be a small step 125 in the wall which continues vertically downward 20 as indicated by the dotted line, which is somewhat exaggerated in Figure 5.

The barrel is flexible and is spread outwardly a slight amount to the position of Figure 6 just prior to retraction. Here the mating surfaces 126, 128 are separated an amount which

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reduces the clamping force on the needle holder 94. The spreading shown in Figure 6 is greatly exaggerated for illustration. It is estimated that an expansion of only about four thousandths of an inch is sufficient to release needle holder 94 from nose 88. By slight further depression of the plunger from the position of Figure 6 to the retracted position of Figure 7, retraction occurs when the retraction force applied by spring 96 exceeds the remaining holding force on needle holder 94. Needle holder 94 then moves through opening 112 into cavity 108 along with a portion of spring 96. The uncompressed length of spring 96 is designed to provide sufficient backward movement to withdraw an injection needle 28 fixed in front portion 94 and carry dislodged stopper 114 with it. At the same time, cap 42 enters opening 138 at the rear of a barrel extension 54 where the peripheral edge is closely confined in order to prevent tampering after retraction.

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The location and configuration of ramp 124 is arranged to avoid cumulation of force required during the retraction sequence. Most of stopper 114 should be dislodged by thumb pressure on plunger 104 before significant resistance develops as angled surfaces 122 begin pushing outwardly on ramp 124. The selection of the location of ramp 24 and the angle of the engaging surfaces make it possible to have a fairly smooth continuous force since the dislodging force continuously decreases as the sliding interface area 116, 118 between the plunger and the stopper is linearly decreased. Because ramp 124 is relatively very small, it is still possible to remove a stepped molding core from the rear of the outer body 84. Alternately, ramp 124 can be the smaller diameter step 125 which avoids reentrant angles whereby resistance to removal of the molding core could occur. After retraction, the back of the plunger is unaccessible and there is no way to reach to stopper or the needle holder in order to reinstall them for re-use.

When used normally, syringe 10 may have a small amount of fluid remaining in the variable chamber in the second position shown in Figure 2 which is, of course, greatly exaggerated in scale. This may amount to no more than a drop or a few drops of fluid in the remaining space above the retraction mechanism. When syringe 10 is fired by pushing down 5 on end cap 48, to the position of Figure 3, the expanding spring and rearwardly moving needle holder carry any remaining fluid up into retraction cavity 38. Surface tension effects hold the tiny droplets in place along the walls of the plunger and no fluid escapes from nose 16. The syringe is normally used to withdraw fluid from a vial. The fluid is injected into a patient followed by immediate retraction of the needle holder and needle in one step. No leakage of fluid from the nose is observed when the syringe is used to inject fluid into a patient.

It has been discovered, however, that if the needle is forcibly prevented from retracting after syringe 10 is "fired" by pushing down until plunger 48 enters opening 58, the small amount of retained fluid from variable chamber 68 can flow into the nose in the space between the needle holder and nose. If the seal around the head of the needle holder is removed while the needle holder is being restrained from retracting, remaining fluid has time to move down into the nose, but it does not leak out from the opening in the front of the nose. Then if the needle holder is suddenly released and allowed to retract normally, it has been found that leakage of fluid from the opening in the front of the nose could be observed. This undesirable scenario was found to occur under the following circumstances. If the syringe is used to draw blood from 20 the patient, the blood filled syringe is removed from the patient and the needle passed through a rubber septum in a sterile vial. The plunger is then depressed to discharge the patient's blood

into the vial. Users expect to depress the plunger fully after the fluid is discharged to retract the needle. When the plunger is depressed fully to cause retraction, the needle cannot retract normally due to the fact it is frictionally held by the rubber septum of the vial. When the empty syringe is then withdrawn from the vial by pulling the needle out of the septum, it immediately 5 retracts. Droplets of fluid were observed on the vial as soon as retraction took place.

Surprisingly, it was found that a small "puff" of air is the source of this problem. If the needle or needle holder is temporarily restrained and prevented from retracting in the normal manner, a brief puff of forwardly directed air is generated when the needle holder is finally allowed to retract. This puff of air was found to emerge from the front of the syringe causing 10 retained fluid trapped around the needle holder to be blown out of the opening left in the nose when the needle holder retracts. It was discovered that if the hollow interior of the plunger is vented, preferably in the area of thumb cap, this condition does not occur and the fluid is entirely retained within the syringe body.

Figures 9 through 16 illustrate the syringe generally designated as syringe 10 with a modification on the end cap or thumb cap on the plunger to provide for venting of the hollow interior of the plunger which is the retraction cavity. Insofar as possible the original numbering 15 of Figures 1 - 4 is retained with primes used to indicate differences.

Head 34' of plunger 32' is preferably slightly modified from plunger head 34 of Figure 2 in the following respects. The elongated plunger has a longitudinally extending generally 20 tubular wall 140 defining a hollow interior along the length of the plunger. The plunger has a head end 34' in front and a rear end portion 142 with a thumb cap 48' behind. The outer side of wall 140 at head end 34' is sealingly surrounded with a resilient plunger seal member 36'

which is like a band with a pair of separated raised rings 144. Plunger seal 36' fits in a depression in the outer surface of wall 140 where it is securely held in position and prevented from longitudinal movement. Seal member 36' is adapted to slide in sealed contact with a tubular wall when the plunger is moved within syringe barrel 14. It is within contemplation of
5 the invention to have a raised piston molded as part of the plastic plunger to serve as a plunger seal in place of a separate rubber plunger seal 36', although the rubber seal member is preferred.

Wall 140 at head end 34' of the plunger 32' has a reduced diameter front portion extending forward from seal member 36' terminating at tip 40 at the front of plunger 32'. Tip 40 defines the opening 41 which leads into the hollow interior 38. The internal structure is as shown in Figure 1. The wall 140 behind tip 40 has a stepped inner side surface comprising a land having an inwardly facing surface and a larger diameter portion extending behind the land into the hollow interior. A separate dislodgeable stopper 42 is slidably held within the reduced diameter front portion of plunger head 34' by a holding force in excess of the fluid injection pressure force to be expected during use of the plunger in syringe barrel 14. Stopper 42 has a back end portion comprising a land 46 and a reduced diameter front end portion extending forwardly beyond tip 40 a fixed distance to its front 146. The fixed distance is the distance between front 146 and tip 40.
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As is seen in Figure 1, the outwardly facing surface 46 of dislodgeable stopper 42 is in sliding sealed engagement with the inwardly facing surface of land 44 in the plunger wall. These lands cooperate to apply a holding force to the stopper and seal hollow interior 38 of
20 plunger 32' from the expected amount of fluid injection pressure force generated in the variable chamber 68 during an injection. The ratio of the effective area of variable chamber 68 to the

area of stopper 42 exposed to fluid pressure is at least two to one and preferably three to one or more as previously indicated. This makes it possible to utilize lower holding forces without blowing out the stopper during an injection. The cooperating lands on the inside of the plunger head and the stopper have sufficient longitudinal length to allow dislodgeable stopper 42 to move 5 the fixed distance between its initial extension at 146 and tip 40 in sliding response to forward movement of the plunger after front 146 of stopper 42 contacts a stop.

As indicated in Figures 1 - 3, front 146 of the stopper 42 encounters head 72 of needle holder 22 which serves as a stop. The fluid opening in head 72 of needle holder 22 is preferably provided with some fine slots or grooves so that fluid can continually enter fluid path 70 as the plunger moves from the position of Figure 1 to that of Figure 2. As the position of Figure 2 10 is reached, the holding force on stopper 42 is reduced by substantial disengagement of the cooperating lands 44, 46 in preparation for dislodgement of the stopper, without unsealing the hollow interior/retraction chamber 38 within plunger 32'. A notch 148 is preferably provided in the tip to prevent trapping fluid at the tip.

Thumb cap 48' at the rear end portion 142 of plunger 32' includes one or more channels 150 which receive vented air from hollow interior 38. Thumb cap 48' has an opening 152 for a closure 154 best seen in Figures 10 and 11. Channels 150 are open at the top for ease of molding although closed channels could also be used.

Figure 10 shows an enlarged top plan view illustrating the use of three channels 150 in 20 combination with a preferred closure 154 installed in circular opening 152. Figure 11 best shows how the channels 150 receive vented air from hollow interior 38. Closure 154 preferably has a stepped outer surface comprising a rear step 156 which rests in opening 152, an

intermediate step 158 which rests in an enlarged portion 160 of the inner side of wall 140 and a front step 162 which rests against inner surface 164 of wall 140. In effect, these structures provide convenient seating for closure 154. Steps 158 and 162 are conveniently provided in a downwardly depending skirt 166.

Importantly, inner surface 164 everywhere there is a channel 150, is provided with a longitudinally extending groove 168 in fluid communication with the hollow interior 38 and the channels 150. Any convenient number may be chosen as the channels are easily molded into the end cap when it is formed. The longitudinally extending grooves 168 do not extend through the entirety of the wall 140 although they could. They are designed for ease of molding since they can be formed in the mold that makes the plunger without using separate pins to form an opening. This is an important cost consideration in a multiple out high speed molding process. This structure is designed for preventing the user's thumb from obstructing the vent opening leading from the interior of the plunger thereby assuring that venting will take place.

Referring now to Figures 9 and 12, it will be noted that opening 58 in the back end of barrel 14 includes slots 172 in fluid communication with the hollow interior of the plunger through one or more channels 150 so that when thumb cap 48' is received in opening 58, no seal is created by the thumb being in contact with opening 58 which might otherwise prevent air from venting. The outer periphery of thumb cap 48' is closely received in opening 58 as the syringe is fired, to prevent reuse. Thumb cap 48' is preferably sized in relation to barrel 14 such that opening 58 is simply an extension in a linear direction of the wall of barrel 14 rather than enlarged as shown. Finally, the interior surface 164 preferably has several annular constrictions 170 designed to catch the head of stopper 42 during its rearward travel. Since stopper 42 is

preferably installed from the rear of the plunger before closure 154 is put in place, the constrictions 170 must allow stopper 42 to be forced through to the front.

A first alternative thumb cap and closure arrangement is illustrated in Figures 13 and 14.

In this embodiment, four channels 150 are provided in thumb cap 48''. Closure 174 has four flat side portions 176 spaced around the periphery at 90° intervals, each in fluid communication with a channel 150. A gap is created at each flat side between the flat sides 176 and the opening 152' which are in fluid communication with interior 38 to create a flow passage for air from interior 38 through the gap along the flat side then into channel 150. Annular groove 178 in closure 174 may be used to fluidly connect each of the flat areas 176 at the level of channels 150. In addition to equalizing air flow, the annular groove allows venting of air regardless of the angular orientation of closure 174 with respect to thumb cap 48''.

A second alternate embodiment has the same thumb cap 48'' with a modified closure 180. Closure 180 has a head 182 which snugly fits within opening 152' which is at the back of the plunger. Opening 152' is only slightly larger than the interior of the plunger to provide a seat for the closure. Four undercut portions 186 are each in joint fluid communication with the interior 38 and one of the channels 150 to create a flow passage from the interior 38. Closure 180 effectively seals the opening 152' so that no fluid particles can escape from the opening. As in the previous embodiment, an annular groove 178 bridges each undercut portion opening into a corresponding channel 150 thereby tying the undercut portions together in fluid communication regardless of the angular orientation of the parts.

In operation, there are many advantages to the improved combination disclosed herein. The diameter of the stopper in both embodiments and the slidable retaining ring member in the

first embodiment, in relation to the diameter across the fluid chamber, makes it possible to produce a syringe which withstands high blowout pressure. By minimizing the effective surface area exposed to the pressurized fluid during an injection, the syringe will withstand injection thumb force of around fifteen to eighteen pounds during injection and at the same time retract 5 in response to as little as five to six pounds of force on the plunger once the injection fluid has been injected. Once the fluid has been injected, cumulation of force required to concurrently operate the retraction mechanism is avoided. First the stopper is moved back and then the needle holder is released. By constricting the diameter of the syringe near a transition zone where the nose begins, a constriction enables the needle holder to be smaller which in turn allows it to fit in a smaller opening with a smaller stopper in the retraction cavity of the hollow 10 plunger.

A vacuum must be pulled in order to fill the syringe. The ring member or the needle holder, as the case may be, must seal the front nose of the syringe body because otherwise vacuum could be lost and fluid could enter the spring area and leak out the front. The hollow outer body and syringe plunger are preferably made from conventional plastic material used for 15 syringes, which has some flexibility. The tolerances on the diameter of mating facing surfaces between the head of the needle holder and the barrel and between the stopper and head of the plunger are not critical in order to maintain a consistent holding and dislodging force. This is believed to be because increasing interference fit increases the frictional holding force only up 20 to a point and then the surrounding wall simply expands a small amount or the internal parts are compressed a small amount without a corresponding increase in the longitudinal force required to move the retainer member or plug member in the retraction direction. It is a desirable self

correcting mechanism which is a cost and quality benefit in making the parts. It is believed that a plastic retainer member could be used and the same self limiting frictional holding force would be obtained.

In the best mode the stopper and the ring member are preferably made from a thermoplastic rubber material designated number 181-55 available from Advanced Elastomer Systems, 540 Maryville Centra Drive, St. Louis, Missouri and sold under the trade name Santoprene®. It is said to have a characteristic hardness around 55 on the Shore A durometer scale which allows for the right amount of resistance to compression, fluid resistance such that the material does not swell when in contact with most fluids, environmental stability allowing the friction and sealing properties to remain non-temperature sensitive, good property retention after aging and excellent property retention after sterilization by all accepted methods. The plunger seal around the head of the plunger is conventional.

The parts are few in number and easily mass produced. The alternate embodiment has the fewest number of separate parts of any tamperproof retractable syringe. The plunger has a one piece hollow outer body with a transition zone and a narrow nose portion. The internal diameter is stepped to greater diameters from front to back for molding around a non-collapsible core which can be extracted from the rear. The same is true for the plunger.

Assembly is greatly simplified and can be accomplished with high speed mechanized equipment. The needle holder and spring are installable from the rear of the barrel without the needle. In the first embodiment the retainer member is forced fit over the inner head of the needle holder and the assembly together with the uncompressed spring are pushed forward and held by sliding engagement of the cooperating inwardly and outwardly facing surfaces while

compressing the spring. The front of the needle holder passes through an opening in the nose which makes it easy to install the needle from the front by conventional means. The alternate embodiment is installed the same way except that there is no separable retainer member around the head of the needle holder.

5 The narrow nose provides a particular advantage for mechanized assembly. The nose has a wall defining an elongated internal cavity which closely confines the spring and needle holder combination. During installation this cavity serves as a guide to steer the needle holder and uncompressed spring into a compressed state of the spring. This solves an important assembly problem. If there is much lateral space in the nose around the spring, when the uncompressed spring is being compressed, it is a laterally unstable column which flexes sideways and bunches up causing a jam up. It might be added that rounded edges on the bottom of the slot directly below retainer 66 would further facilitate entry of the end of the spring.

10 The stopper is also installable from the rear of the plunger by pushing it forward until the cooperating lands are slidably engaged. Then plug member 50 is force fit or otherwise fixed in the opening at the back of the plunger and the plunger is installed in the outer body. It is not necessary to try to pass the sharp needle through an elongated body with constricted openings where slight misalignment could cause hangups. The head of the needle holder simultaneously acts as a seal as well as a holding device such that no seal is required at the tip of the nose and no ultrasonic welding of separate parts is required.

15 There is no necessity for using internal locking teeth of any kind. No locking teeth are needed to hold the retraction mechanism or to lock the plunger after retraction. Locking teeth present difficult molding and quality control problems, tend to be temperature sensitive and tend

to require a larger diameter barrel which increases premature blowout problems. In addition to
the non-reusability provided by separation of the retainer ring from the head of the needle holder
and dislodgement of the stopper, the plunger is not accessible after retraction because it is
depressed within an opening at the back of the outer body. This additional tamperproof feature
5 is provided in a one piece body without the necessity for hooking anything or twisting anything.
The easily made and installed force fit plug at the back of the retraction cavity prevents access
to the retracted components. The Federal government has rights in the invention under 35
U.S.C. §203. The Federal government has a nonexclusive, nontransferable irrevocable, paid
up license to the invention as set forth in the priority documents.

I claim:

1. A syringe plunger handle assembly for use in a syringe barrel to inject fluid, comprising:

an elongated plunger formed with a longitudinally extending generally tubular wall defining a hollow interior along the length of the plunger, the plunger having a head end in front and a rear end portion with a thumb cap behind;

the outer side of the wall at the head end has a plunger seal adapted to slide in sealed contact with a tubular wall when the plunger is moved within a syringe barrel;

the wall at the head end of the plunger has a reduced diameter front portion extending forwardly beyond the plunger seal to a tip at the front of the plunger defining an opening within the front end portion leading into the hollow interior, the wall behind the tip having a stepped inner side surface comprising a land having an inwardly facing surface and a larger diameter portion extending behind the land into the hollow interior;

a separate dislodgeable stopper slidingly held within the reduced diameter front portion of the plunger head by a holding force in excess of an expected fluid injection pressure force during use of the plunger in a syringe barrel, the stopper having a back end portion comprising a land and a reduced diameter front end portion extending a fixed distance forwardly beyond the tip, the stopper land having an outwardly facing surface in sliding sealed engagement with the inwardly facing surface of the land in the plunger wall, said lands comprising cooperating lands which apply said holding force to the stopper and seal the hollow interior of the plunger from said expected fluid injection pressure force; and

the cooperating lands have sufficient length to allow the dislodgeable stopper to move said fixed distance back to the tip in sliding response to forward movement of the plunger after the front of the stopper contacts a stop, whereby said holding force is reduced by substantial disengagement of the cooperating lands in preparation for dislodgement of the stopper without unsealing the hollow interior of the plunger.

2. The syringe plunger handle assembly of Claim 1 wherein a vent for the hollow interior is located at the rear end portion of the plunger.

3. The syringe plunger handle assembly of Claim 2 wherein the thumb cap at the rear end portion of the plunger is provided with a channel which receives vented air from the hollow interior.

4. The syringe plunger handle assembly of Claim 3 wherein said vent comprises at least one groove at the rear end portion of the plunger in fluid communication with said channel for venting the retraction cavity in the plunger.

5. The syringe plunger handle assembly of Claim 1 wherein the thumb cap at the rear end portion of the plunger surrounds an opening suitable for installation of the dislodgeable stopper into the hollow interior and a closure for the opening in the thumb cap.

6. The syringe plunger handle assembly of Claim 5 wherein at least one channel in the thumb cap receives vented air from the hollow interior.
7. The syringe plunger handle assembly of Claim 6 wherein the closure has at least one cutaway side portion in joint fluid communication with the hollow interior and said channel to create a vent passage.
8. The syringe plunger handle assembly of Claim 7 wherein the rear end portion of the plunger has an internal surface with at least one groove in fluid communication with the hollow interior and said at least one channel in the thumb cap.
9. The syringe plunger handle assembly of Claim 5 wherein the closure has a headed portion fitted in the opening in the thumb cap and a skirted side containing an undercut configured to reside in fluid communication jointly with the hollow interior and at least one channel in the thumb cap to comprise at least one vent passage from the hollow interior.

10. A syringe plunger handle assembly and syringe barrel combination for use in injecting fluids, comprising:

a hollow syringe body having a longitudinally extending tubular wall comprising an elongated barrel portion having a back end portion with an opening behind and a nose in front, 5 said nose having a reduced cross-sectional area relative to the barrel;

an elongated plunger formed with a longitudinally extending generally tubular wall defining a hollow interior along the length of the plunger, the plunger having a head end in front mounted in the syringe body and a rear end portion with a thumb cap behind extending from said opening in the barrel;

10 the outer side of the wall at the head end has a plunger seal adapted to slide in sealed contact with the tubular wall when the plunger is moved within the syringe barrel;

the wall at the head end of the plunger has a reduced diameter front portion extending forwardly beyond the plunger seal to a tip at the front of the plunger defining an opening within the front end portion leading into the hollow interior, the wall behind the tip having a stepped inner side surface comprising a land having an inwardly facing surface and a larger diameter portion extending behind the land into the hollow interior;

20 a separate dislodgeable stopper slidingly held within the reduced diameter front portion of the plunger head by a holding force in excess of an expected fluid injection pressure force during use of the plunger in the syringe barrel, the stopper having a back end portion comprising a land and a reduced diameter front end portion extending a fixed distance forwardly beyond the tip, the stopper land having an outwardly facing surface in sliding sealed engagement with the inwardly facing surface of the land in the plunger wall, said lands comprising cooperating lands

which apply said holding force to the stopper and seal the hollow interior of the plunger from expected fluid injection pressure force;

the cooperating lands have sufficient length to allow the dislodgeable stopper to move said fixed distance back to the tip in sliding response to forward movement of the plunger after 5 the front of the stopper contacts a stop, whereby said holding force is reduced by substantial disengagement of the cooperating lands in preparation for dislodgement of the stopper without unsealing the hollow interior of the plunger; and

further forward movement of the plunger causing dislodgement of the stopper form the opening accompanied by reception of the thumb cap within the confines of the opening at the back end portion of the barrel thereby preventing reuse.

11. The combination of Claim 10 wherein the hollow interior of the plunger is provided with a vent passage in a rear portion thereof behind the plunger seal member.

12. The combination of Claim 11 wherein the thumb cap at the rear end portion of the plunger is provided with a channel which receives vented air form the hollow interior.

13. The combination of Claim 12 wherein the back end portion of the barrel of the syringe body includes at least one slot which allows vented air to escape without being trapped by a user's thumb during retraction when the thumb cap is pressed down and received in said opening in the barrel at the back of the syringe body.

14. The combination of Claim 12 wherein said vent comprises at least one groove at the rear end portion of the plunger in fluid communication with said channel for venting the retraction cavity in the plunger.

15. The combination of Claim 14 wherein the thumb cap at the rear end portion of the plunger surrounds an opening suitable for installation of the dislodgeable stopper into the hollow interior and a closure for the opening in the thumb cap.

16. The combination of Claim 15 wherein the closure has at least one cutaway side portion in joint fluid communication with the hollow interior and said channel to create a vent passage.

17. The combination of Claim 16 wherein the rear end portion of the plunger has an internal surface with at least one groove in fluid communication with the hollow interior and said at least one channel in the thumb cap.

18. The combination of Claim 17 wherein the back end portion of the barrel of the syringe body includes at least one slot which allows vented air to escape without being trapped by a user's thumb during retraction when the thumb cap is pressed down and received in said opening in the barrel at the back of the syringe body.

19. A tamperproof retractable non-reusable syringe structure designed to have a high blowout pressure coupled with a low retraction force, comprising:

a hollow outer body having a longitudinally extending wall comprising an elongated barrel having a back end portion with an opening behind and a nose in front, said nose having
5 a reduced cross-sectional area relative to the barrel;

a retraction mechanism having retractable parts comprising an elongated needle holder and spring combination mounted in the nose portion of the barrel with the needle holder temporarily held in a use position with a needle attached to the needle holder extending from the nose of the barrel and a fluid path through the needle and needle holder leading to a variable fluid chamber in the barrel;
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a hollow plunger handle assembly disposed partially within the elongated barrel of the outer body, the plunger having a head in front with a piston fixedly mounted thereon in slidable sealed contact with the interior of the barrel, a back end portion having a graspable end cap and a retraction cavity therein, the plunger head having a tip in front defining an opening into the retraction cavity and a separate dislodgeable stopper slidingly sealing the opening, said stopper having a front end portion extending a distance beyond the tip, the stopper being capable of sliding relative to the plunger back to the tip without unsealing the opening and without completely dislodging in response to forward motion of the plunger after the front end portion of the stopper has come into contact with the needle holder;
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20 the combined head of the plunger and the stopper defining the upper boundary of the variable fluid chamber;

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the plunger being movable to a second position while expelling fluid contained in the variable chamber through the fluid path in response to thumb pressure on said cap, said second position comprising the end of an injection cycle including sliding of the stopper said distance relative to the plunger tip after contact with the needle holder at the end of the injection cycle before retraction occurs;

the plunger being further depressible to a retraction position beyond said second position whereby the retraction mechanism temporarily holding the needle holder releases the needle holder to carry said stopper and said needle into the retraction cavity and said graspable end cap is simultaneously recessed into the opening at the back of the barrel so the syringe cannot be reused.

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20. The tamperproof retractable non-reusable syringe structure of Claim 19 wherein the plunger has at least one vent passage spaced behind the piston, in fluid communication with the retraction cavity, to allow venting of air from the retraction cavity in response to entry of retractable parts therein, sufficient to prevent compression and blow by of air into the variable fluid chamber which could otherwise expel residual fluid from the nose of the syringe.

21. The tamperproof retractable non-reusable syringe structure of Claim 20 wherein the thumb cap at the rear end portion of the plunger surrounds an opening suitable for installation of the dislodgeable stopper into the plunger and a closure for the opening.

22. The tamperproof retractable non-reusable syringe structure of Claim 21 wherein said at least one vent passage comprises a channel in the thumb cap leading outward from the opening at the rear end portion of the plunger.

23. The tamperproof retractable non-reusable syringe structure of Claim 22 wherein the closure has at least one cutaway side portion in fluid communication with the retraction cavity and said channel to create said at least one vent passage.

24. The tamperproof retractable non-reusable syringe structure of Claim 23 wherein the wall of the barrel near the opening includes at least one slot which allows vented air to escape without being trapped by a user's thumb during retraction when the thumb cap is pressed down, received and recessed in the opening in the back end portion of the barrel of the syringe body.

25. The tamperproof retractable non-reusable syringe structure of Claim 22 wherein the back end portion of the plunger has an internal surface with at least one groove in fluid communication with said channel in the thumb cap to comprise said at least one vent passage.

26. The tamperproof retractable non-reusable syringe structure of Claim 25 wherein the back end portion of the barrel of the syringe body includes at least one slot which allows vented air to escape without being trapped by users thumb during retraction when the thumb cap is pressed down and received in the opening of the barrel at the back of the syringe body.

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27. The tamperproof retractable non-reusable syringe structure of Claim 21 wherein the closure has a headed portion fitted in the opening at the rear end of the plunger and a depending skirted side containing an undercut configured to reside in fluid communication jointly with the retraction cavity and at least one channel in the thumb cap to comprise said at least one vent passage.

28. The tamperproof retractable non-reusable syringe structure of Claim 27 wherein the back end portion of the barrel of the syringe body includes at least one slot which allows vented air to escape without being trapped by a user's thumb during retraction when the thumb cap is pressed down and received in said opening in the barrel at the back of the syringe body.

ABSTRACT

A tamperproof retractable non-reusable syringe has a one piece hollow outer body with a barrel for a slidable plunger, a transition zone and a smaller diameter nose portion. An elongated needle holder and spring combination is installable from the rear of the outer body, guided into the nose portion and held by cooperating inwardly and outwardly facing surfaces oriented in the direction of retraction at the most constricted part of the transition zone where the nose begins. The plunger has an opening with a dislodgable stopper for receiving parts of the retraction mechanism. The stopper and the head of the needle holder are of significantly reduced diameter from the injection fluid chamber to resist blowing out prematurely. In one embodiment the head of the needle holder is surrounded by a separable retainer member which is slidingly removed by contact with the tip of the plunger after the stopper is mostly or fully removed to avoid cumulation of force required for retraction after the injection. In a second embodiment the head of the needle holder is clamped and held by constricting forces imposed by stress on the outer body induced by interference fit. Release occurs by slight expansion on the barrel by contact of the plunger tip with a small internal ramp in the outer barrel. Both embodiments have a plunger cap configured to enter an opening in the outer body to provide an additional tamperproof feature. The retraction cavity is provided with venting structures to assure that all uninjected fluid is retained within the syringe body.

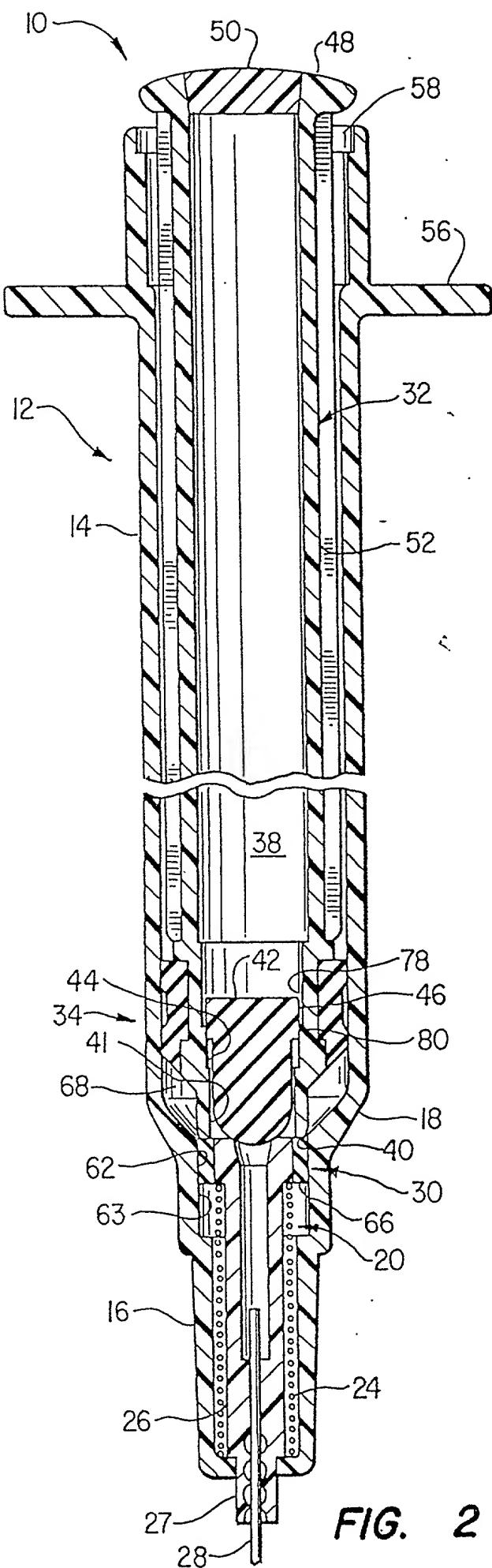
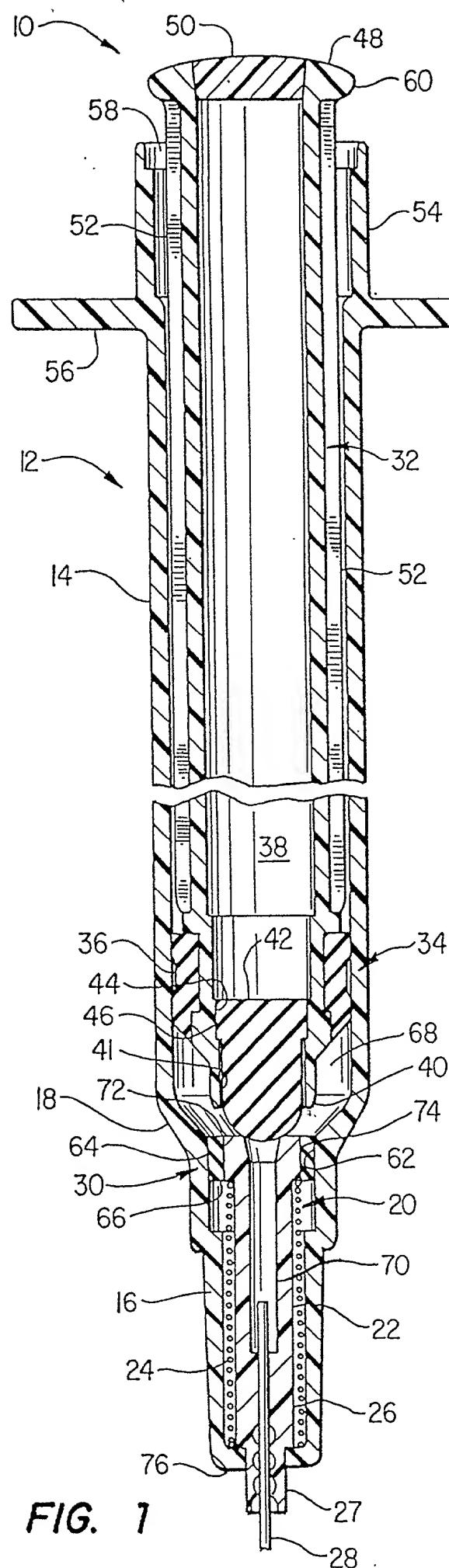
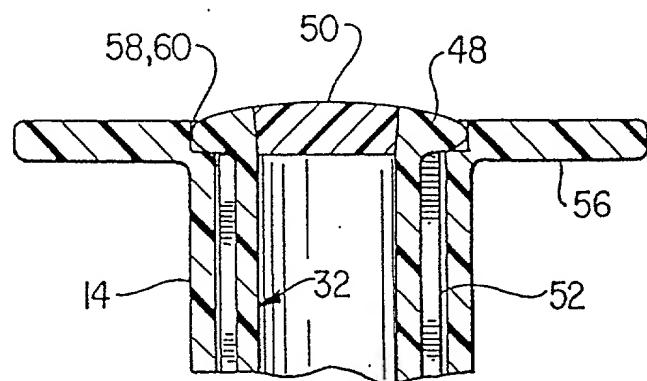
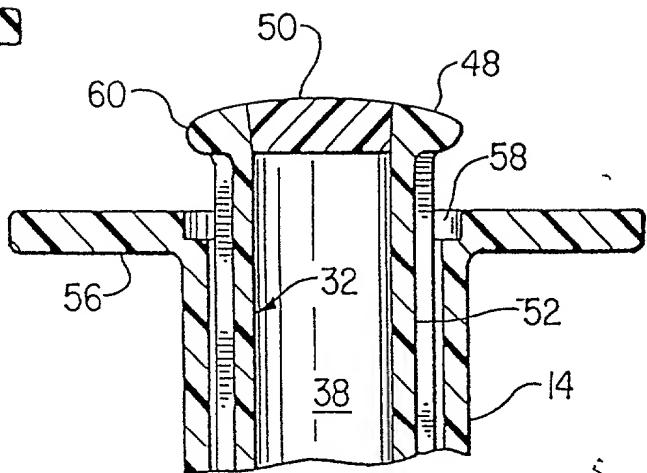
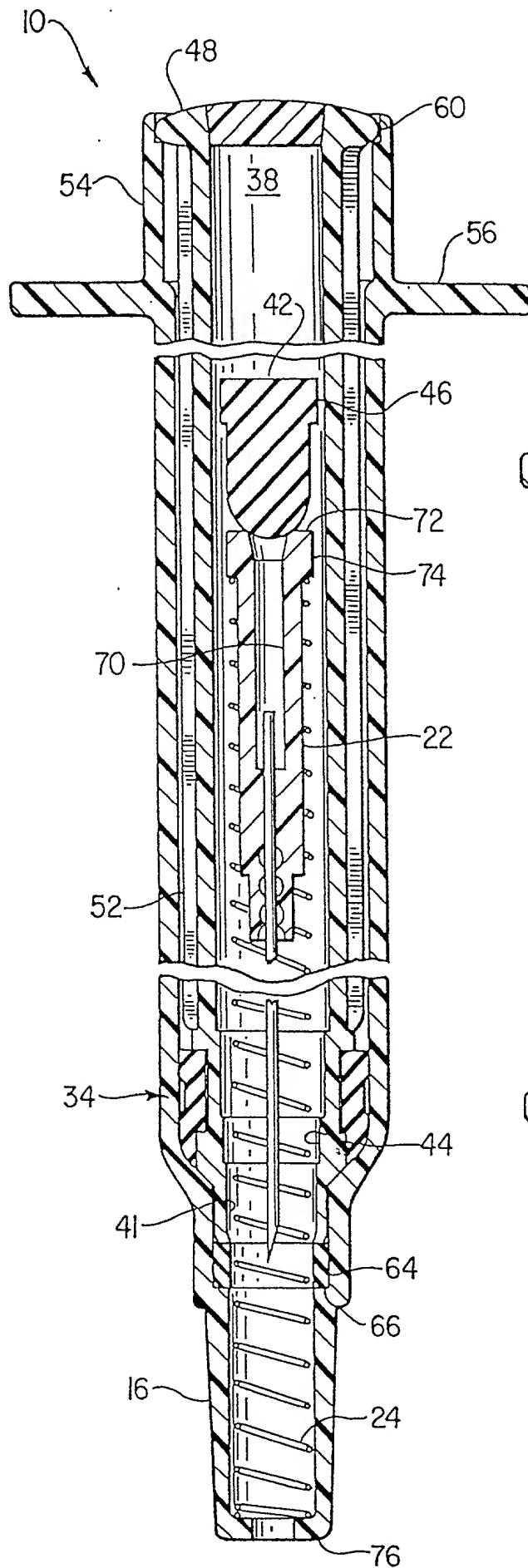
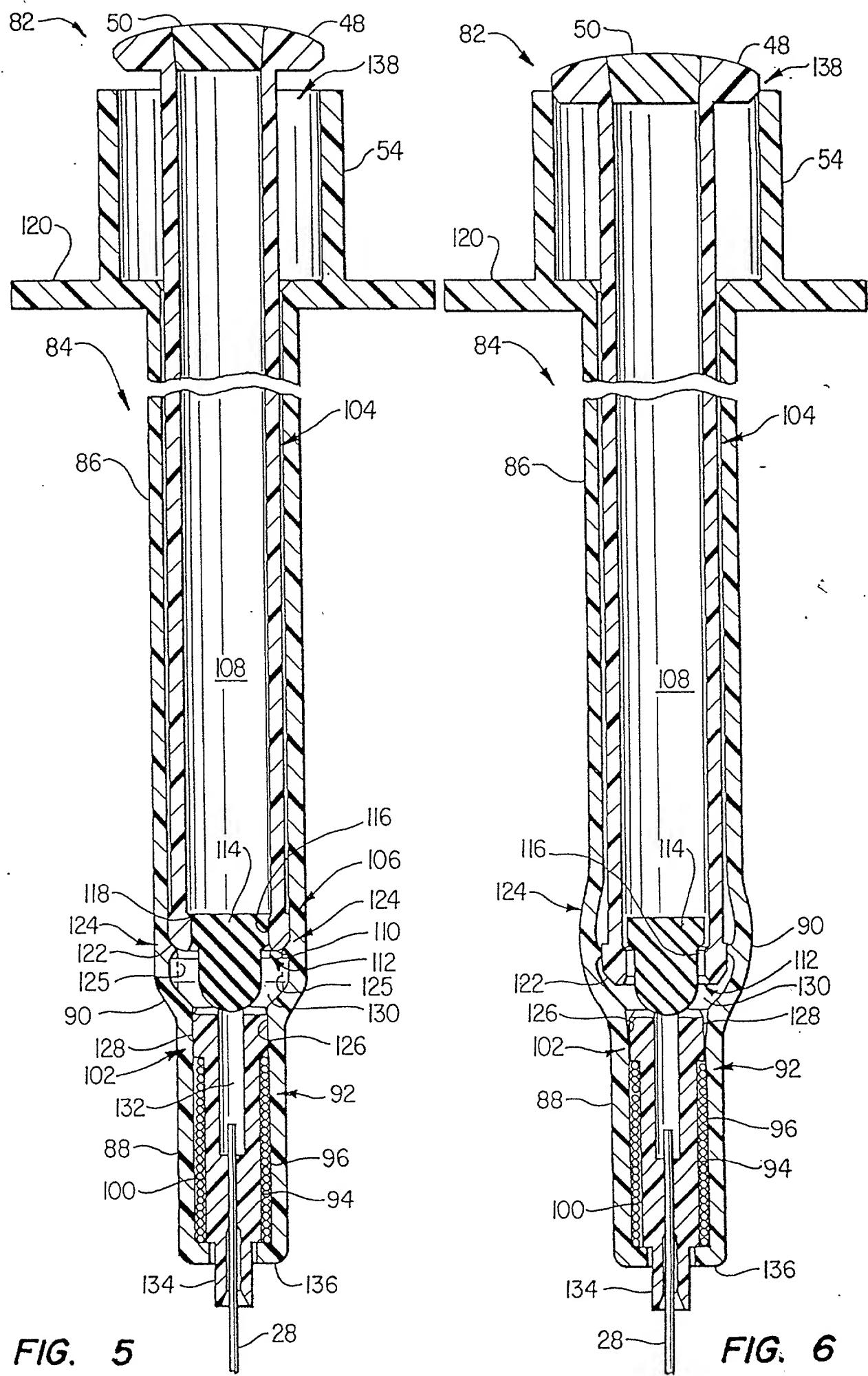


FIG. 1

FIG. 2





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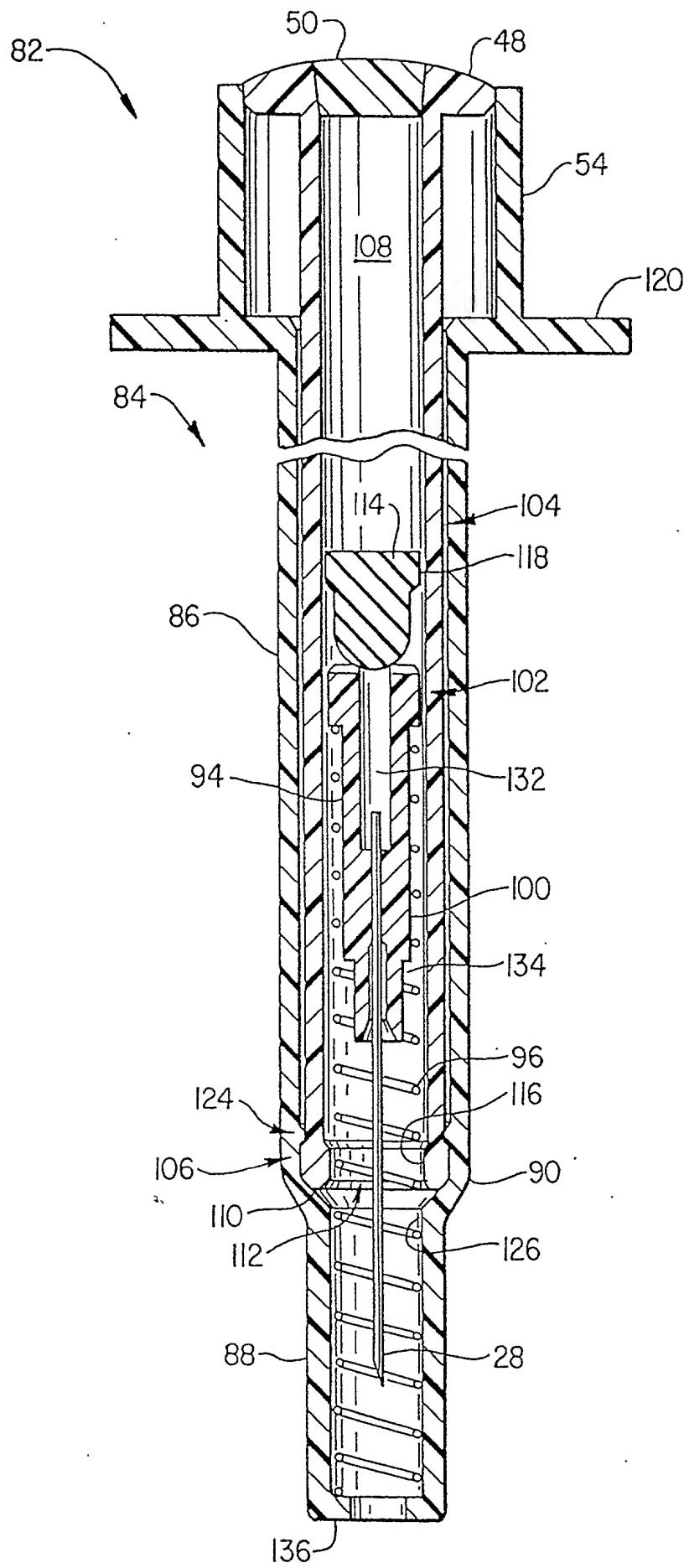


FIG. 7

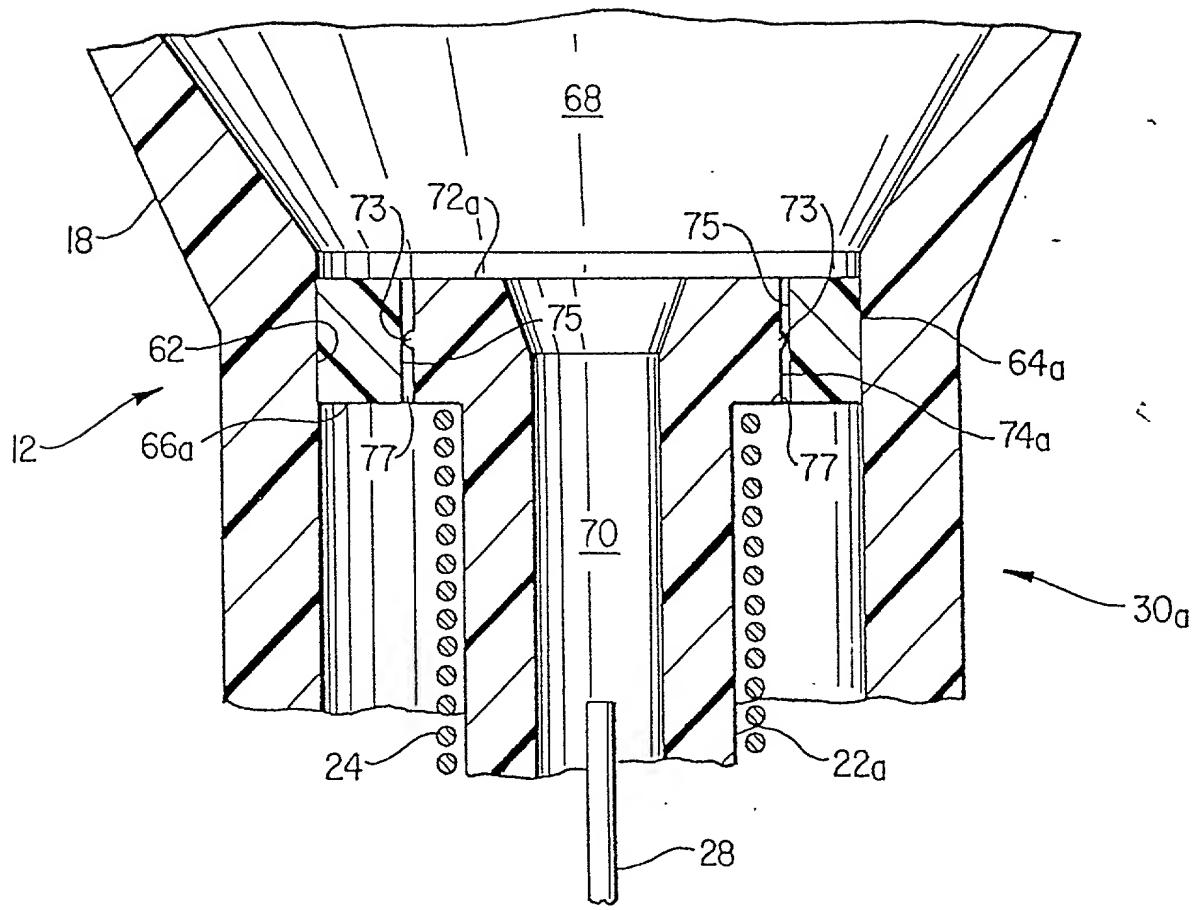
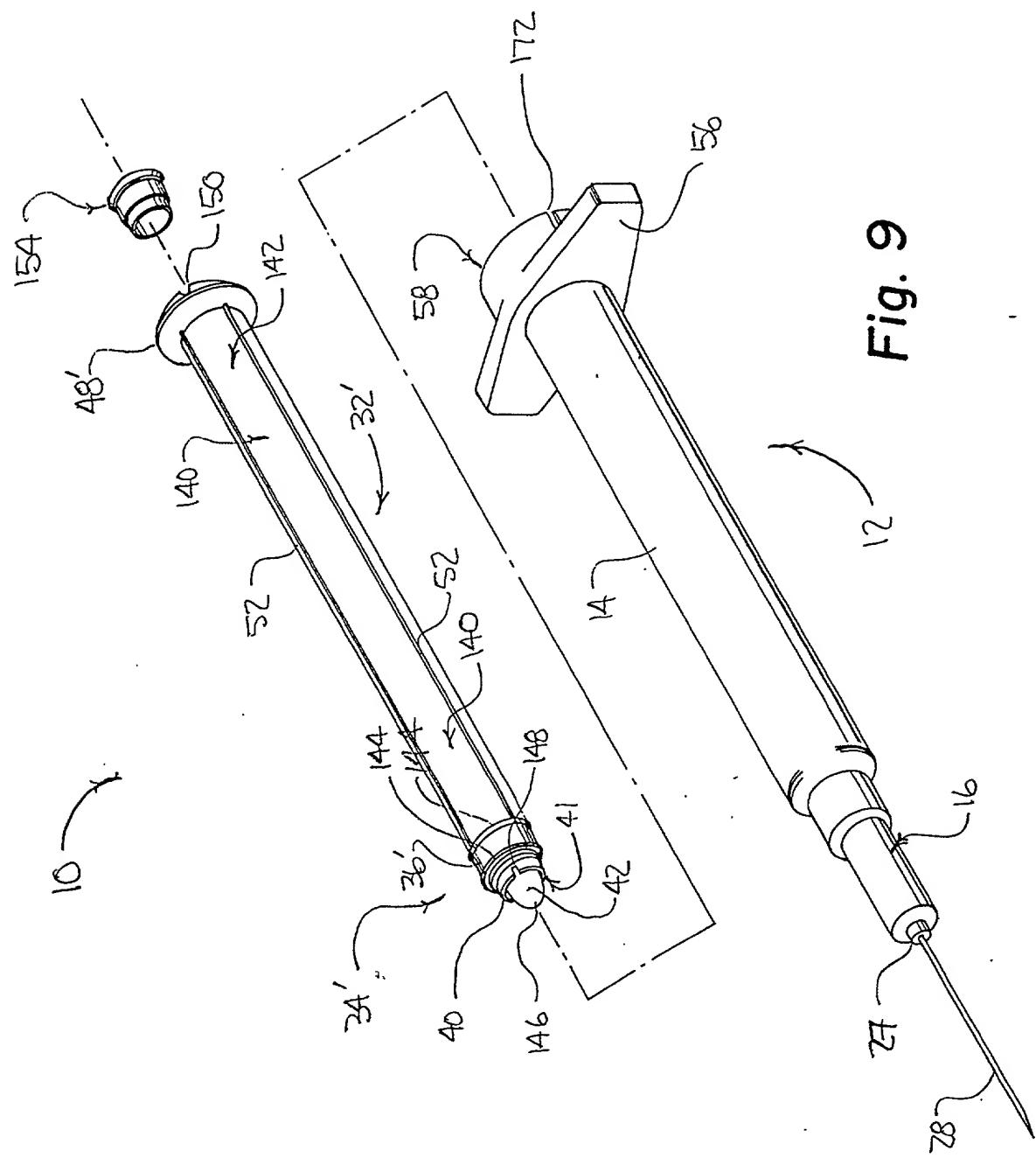


FIG. 8

Fig. 9



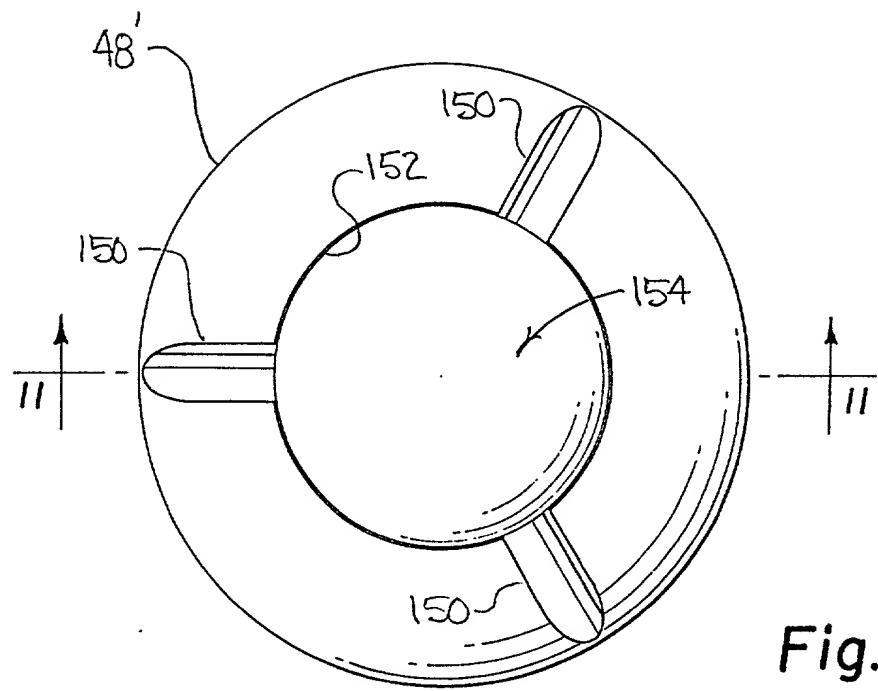


Fig. 10

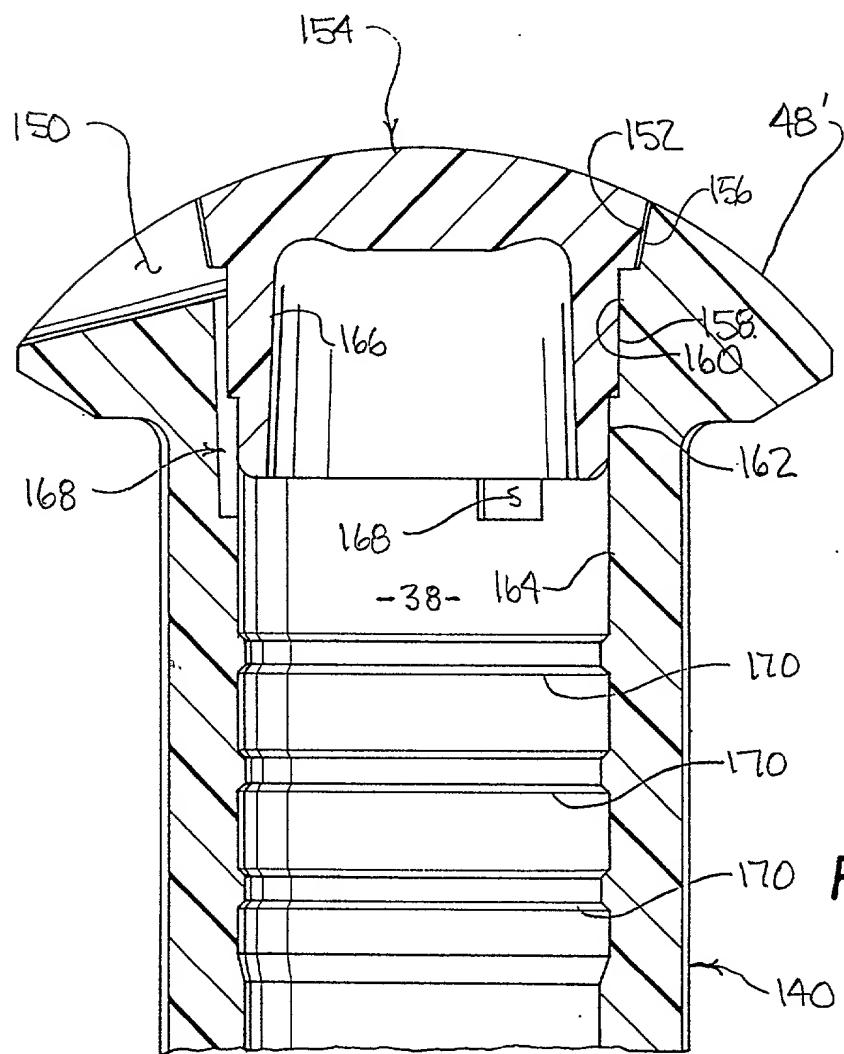


Fig. 11

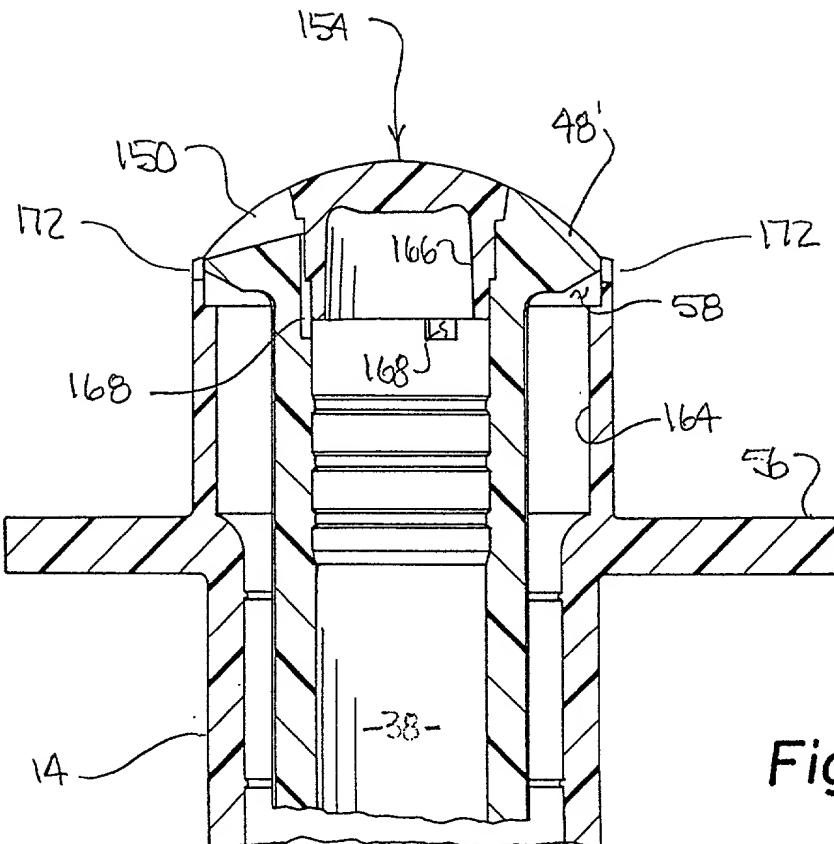


Fig. 12

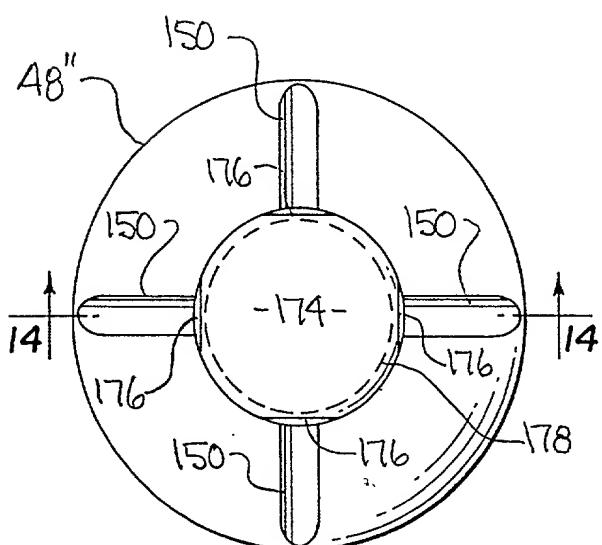


Fig. 13

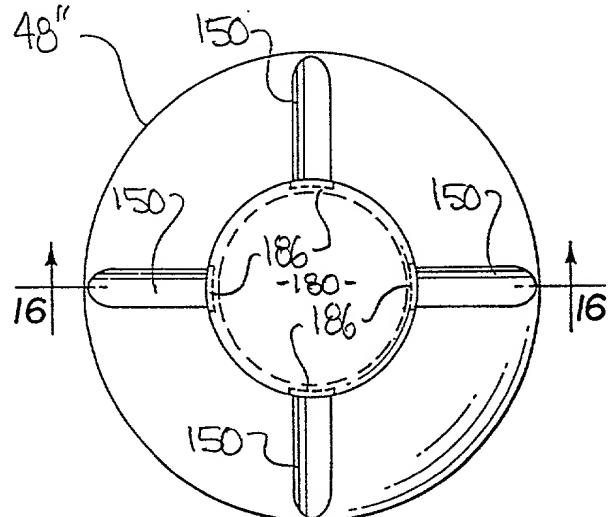


Fig. 15

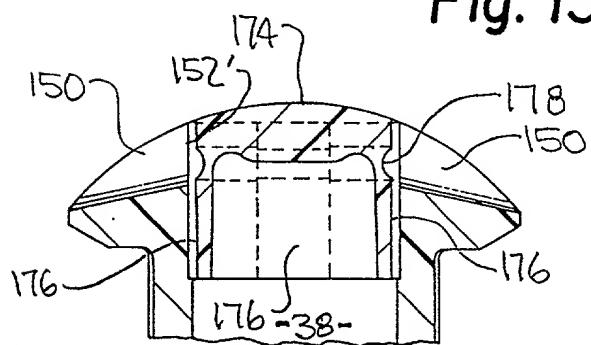


Fig. 14

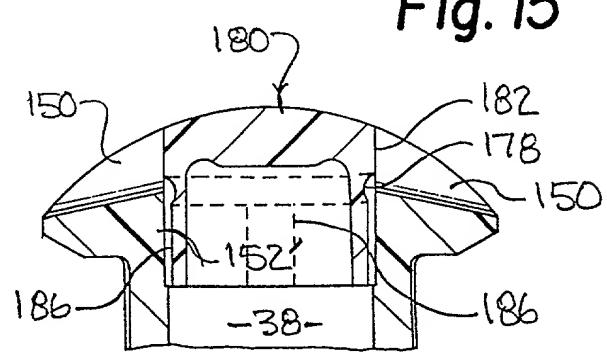


Fig. 16

DECLARATION AND POWER OF ATTORNEY

As the below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled: "SYRINGE PLUNGER ASSEMBLY AND BARREL".

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to herein.

I acknowledge the duty to disclose to the Office all information known to me to be material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

NONE

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Section 112, I acknowledge the duty to disclose to the Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

Serial Number 08/537,242, filed September 29, 1995 and Serial Number 08/438,954, filed May 11, 1995, now U.S. Patent 5,578,011, entitled Tamperproof Retractable Syringe.

And I hereby appoint HARRY J. WATSON, Registration No. 29,985; L. DAN TUCKER, Registration No. 22,670; WILLIAM D. HARRIS, JR., Registration No. 19,243; ROY W. HARDIN, Registration No. 28,304; WILLIAM D. JACKSON, Registration No. 20,846; MICHAEL S. BUSH, Registration No. 31,745; KRISTIN K. JORDAN, Registration No. 37,859; MICHAEL W. PIPER, Registration No. 39,800; and CRAIG J. COX, Registration No. 39,643 all of the firm of HARRIS, TUCKER & HARDIN, P.C., my attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith. I request that all correspondence be addressed to:

Harry J. Watson
HARRIS, TUCKER & HARDIN, P.C.
One Galleria Tower
13355 Noel Road, Suite 2100
Dallas, Texas 75240-6604
972/233-5712

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of inventor: Thomas J. Shaw

Inventor's signature Thomas J. Shaw

Date 4/21/97

Residence: 1510 Hillcrest
Little Elm, Texas 75068

Citizenship: United States

Post Office Address: 1510 Hillcrest
Little Elm, Texas 75068

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